

TITLE OF THE INVENTION

COCCIDIAN PARASITE CASEIN KINASE I AS A CHEMOTHERAPEUTIC TARGET FOR ANTIPROTOZOAL AGENTS.

5 CROSS-REFERENCE TO RELATED APPLICATIONS

The present application claims the benefit of U.S. Provisional Application No. 60/537,094, filed January 16, 2004, hereby incorporated by reference herein.

10 FIELD OF THE INVENTION

The present invention relates in part to isolated nucleic acid molecules (polynucleotides) which encode coccidian casein kinase I ("CKI") enzymes, members of the family of serine/threonine CKI proteins. Specifically, the present invention relates to three disclosed CKI proteins isolated from the protozoan species *Eimeria tenella* (EtCKI α) and *Toxoplasma gondii* (TgCKI α and TgCKI β). The present invention further relates to recombinant
15 vectors and recombinant hosts which contain a DNA fragment encoding a coccidian CKI protein, substantially purified forms of coccidian CKI proteins, and recombinant membrane fractions and/or cell lysates comprising said proteins. Characterization of the coccidian CKI proteins of the present invention allows for screening methods to identify novel coccidiostat compounds that may have therapeutic activity for veterinary and human diseases, including coccidiosis and
20 Toxoplasmosis. Thus, the present invention also relates to methods associated with identifying compounds which specifically modulate coccidian CKI activity.

BACKGROUND OF THE INVENTION

Protein kinases constitute the largest family of human enzymes. As principal
25 regulators of signal transduction pathways, protein kinases play a critical role in a wide range of cellular processes such as proliferation, differentiation and apoptosis. Protein kinases are currently considered to be the largest class of proteins amenable to therapeutic intervention by small molecule drugs (Cohen, P., 2002, *Nat. Rev. Drug Discov.* 1:309-315; Hopkins, A.L. and Groom, C.R., 2002, *Nat. Rev. Drug Discov.* 1:727-730). With the advent of genome databases of
30 clinically important protozoan parasites, it is clear that the evolutionary distance between essential, protozoan protein kinases and the host organisms' corresponding kinase orthologues presents many opportunities for chemotherapeutic intervention of protozoan parasitic proliferation (Doerig, C. et al., 2002, *Trends Parasitol.* 18:366-371). Such differences have been successfully exploited in the case of the coccidian parasite cGMP-dependent protein kinase
35 ("PKG") for which a selective, small molecule inhibitor has been identified that effectively

controls parasite proliferation, preventing coccidiosis caused by *Eimeria* species in poultry and Toxoplasmosis caused by *Toxoplasma gondii* in a mouse model (Gurnett, A.M. et al., 2002, *J. Biol. Chem.* 277:15913-15922; Nare, B. et al., 2002, *Antimicrob. Agents Chemother.* 46:300-307; Donald, R.G.K. et al., 2002, *Eukaryot. Cell.* 1:317-328). Knockaert et al. (2000, *Chem. Biol.* 7:411-422) also demonstrated the exploitable differences between parasite kinases and their host cell counterparts using affinity chromatography with an immobilized inhibitor of cyclin-dependent protein kinase ("CDK"), purvalanol B. While the ligand used in these studies is an exceptionally potent and selective inhibitor of mammalian CDKs, affinity chromatography of parasite lysates, including *Plasmodium falciparum*, *Leishmania mexicana*, *Toxoplasma gondii* and *Trypanosoma cruzi*, with this inhibitor showed protozoan casein kinase I ("CKI") proteins, rather than CDK enzymes, tightly bound to the column. In contrast, parallel affinity chromatography of extracts of marine invertebrates and some vertebrate tissues positively identified associated CDKs without selection of CKI enzymes.

Casein kinase I enzymes represent a family of multipotential serine/threonine proteins kinases common to all eukaryotic cells. These enzymes are known to play important and diverse roles in vesicular trafficking, DNA repair, cell cycle progression and cytokinesis in organisms from yeast to humans. In multicellular organisms they also regulate developmental pathways, control circadian rhythms, and have been implicated in Alzheimer's disease progression. Seven members of the CKI family have been identified (α , β , $\gamma 1$, $\gamma 2$, $\gamma 3$, δ , and ϵ isoforms). It is likely that CKI isoforms also play an essential role in protozoan parasites since these enzymes have been described for *Plasmodium*, *Leishmania* and *Trypanosoma* parasites (Barik, S. et al., 1997, *J. Biol. Chem.* 272:26132-26138; Vieira, L.L. et al., 2002, *Int. J. Parasitol.* 32:1085-1093; Sacerdoti-Sierra, N. and Jaffe, C.L., 1997, *J. Biol. Chem.* 272:30760-30765; Spadafora, C. et al., 2002, *Mol. Biochem. Parasitol.* 124:23-36; Calabokis, M. et al., 2002, *Parasitol. Int.* 51:25-39). The present invention relates to the cloning, expression and characterization of three novel CKI enzymes, two isoforms from the species *Toxoplasma gondii* (TgCKI α and TgCKI β) and one from the species *Eimeria tenella* (EtCKI α).

Affinity-ligand purification studies from Knockaert et al., 2000, *supra*, using an immobilized purvalanol column identified a single binding protein from *T. gondii* lysates. Microsequencing of an eleven amino acid peptide demonstrated that the *Toxoplasma* protein displayed sequence similarity to *Arabidopsis thaliana* CKI, and thus the protein was labeled *Toxoplasma* CKI.

Eimeria and *Toxoplasma* are related coccidian protozoa, a subgroup of the phylum *Apicomplexa* that includes intestinal parasites of veterinary and clinical significance. The poultry industry is most severely affected by *Eimeria spp.* infections resulting in coccidiosis.

Worldwide costs of \$800 million have been reported by the industry, encompassing the cost of prophylactic in-feed medications, alternative treatments if those medications fail, and losses due to mortality and poor feed conversions of infected birds (Allen, P.C. and Fetterer, R.H., 2002, *Clin. Microbiol. Rev.* 15:58-65). Anticoccidial compounds introduced nearly 30 years ago
5 continue to be used prophylactically in poultry operations, the most successful being the polyether ionophores. However, reports of resistance to the current compounds are common due to the constant chemotherapeutic pressure exerted by this class of compounds. *Toxoplasma gondii* infects a broad range of warm-blooded animals. Although it is usually benign in humans, Toxoplasmosis can result in significant mortality and/or morbidity in congenital infections and
10 immunocompromized patients. Current treatment for Toxoplasmosis is a combination therapy using pyrimethamine and sulfonamide; however, significant toxicity often accompanies this treatment regime. There is a current need for identification and development of new compounds for treatment of Toxoplasmosis and coccidiosis. To meet this end, the coccidian casein kinase I enzymes disclosed herein represent novel targets for the development of broad-spectrum
15 coccidiostat compounds effective against coccidiosis and Toxoplasmosis.

SUMMARY OF THE INVENTION

The present invention relates to an isolated or purified nucleic acid molecule (polynucleotide) which encodes a coccidian casein kinase I ("CKI") protein, a member of the
20 family of multipotential serine/threonine protein kinases common to all eukaryotic cells and known to play important and diverse cellular roles.

The present invention also relates to an isolated or purified nucleic acid molecule (polynucleotide) which encodes a coccidian CKI protein of the *Eimeria* genus, including but not limited to an *Eimeria* CKI protein from the species *Eimeria tenella*.

25 The present invention further relates to an isolated or purified nucleic acid molecule (polynucleotide) which encodes a coccidian CKI protein of the *Toxoplasma* genus, including but not limited to a *Toxoplasma* CKI protein from the species *Toxoplasma gondii*.

The present invention relates to an isolated or purified nucleic acid molecule (polynucleotide) encoding a coccidian CKI protein, this nucleic acid molecule comprising a
30 nucleotide sequence encoding an amino acid sequence selected from the group consisting of SEQ ID NO:2, SEQ ID NO:4 and SEQ ID NO:6.

The present invention also relates to an isolated or purified nucleic acid molecule (polynucleotide) comprising a DNA molecule selected from the group consisting of SEQ ID
NO:1, SEQ ID NO:3 and SEQ ID NO:5, which encode novel coccidian CKI proteins as set forth
35 in SEQ ID NO:2, SEQ ID NO:4 and SEQ ID NO:6, respectively.

The present invention relates to an isolated or purified nucleic acid molecule (polynucleotide) which encodes a coccidian CKI protein, this nucleic acid molecule consisting of a nucleotide sequence encoding an amino acid sequence selected from the group consisting of SEQ ID NO:2, SEQ ID NO:4 and SEQ ID NO:6.

5 The present invention also relates to an isolated or purified nucleic acid molecule (polynucleotide) consisting of a DNA molecule selected from the group consisting of SEQ ID NO:1, SEQ ID NO:3 and SEQ ID NO:5, which encode novel coccidian CKI proteins as set forth in SEQ ID NO:2, SEQ ID NO:4 and SEQ ID NO:6, respectively.

10 A preferred aspect of this portion of the present invention is disclosed in Figure 1, designated as EtCKI α (SEQ ID NO:1), which encodes a novel coccidian CKI protein from the species *Eimeria tenella*.

Another preferred aspect of this portion of the present invention is disclosed in Figure 2, designated as TgCKI α (SEQ ID NO:3), which encodes a novel coccidian CKI protein from the species *Toxoplasma gondii*.

15 Another preferred aspect of this portion of the present invention is disclosed in Figure 3, designated as TgCKI β (SEQ ID NO:5), which encodes a novel coccidian CKI protein from the species *Toxoplasma gondii*.

20 The isolated nucleic acid molecules of the present invention may include a deoxyribonucleic acid molecule (DNA), such as genomic DNA and complementary DNA (cDNA), which may be single (coding or noncoding strand) or double stranded, as well as synthetic DNA, such as a synthesized, single stranded polynucleotide. The isolated nucleic acid molecule of the present invention may also include a ribonucleic acid molecule (RNA).

25 The present invention also relates to biologically active fragments or mutants of SEQ ID NO:1 which encode mRNA expressing a novel *Eimeria tenella* CKI protein. Any such biologically active fragment and/or mutant will encode either a protein or protein fragment which at least substantially mimics the enzymatic properties of the *E. tenella* CKI protein as set forth in SEQ ID NO:2. Any such polynucleotide includes but is not necessarily limited to nucleotide substitutions, deletions, additions, amino-terminal truncations and carboxy-terminal truncations such that these mutations encode mRNA which express a protein or protein fragment of

30 diagnostic, therapeutic or prophylactic use and would be useful for screening for agonists and/or antagonists of CKI activity.

35 The present invention also relates to biologically active fragments or mutants of SEQ ID NOs:3 and 5 which encode mRNA expressing novel *Toxoplasma gondii* CKI proteins. Any such biologically active fragment and/or mutant will encode either a protein or protein fragment which at least substantially mimics the enzymatic properties of the *T. gondii* CKI

protein as set forth in SEQ ID NO:4 and SEQ ID NO:6, respectively. Any such polynucleotide includes but is not necessarily limited to nucleotide substitutions, deletions, additions, amino-terminal truncations and carboxy-terminal truncations such that these mutations encode mRNA which express a protein or protein fragment of diagnostic, therapeutic or prophylactic use and would be useful for screening for agonists and/or antagonists of CKI activity.

The present invention further relates to a purified nucleic acid molecule (polynucleotide) encoding a coccidian CKI protein, wherein said nucleic acid molecule comprises a nucleotide sequence selected from the group consisting of: (a) a nucleotide sequence which encodes an amino acid sequence as set forth in SEQ ID NO:2, SEQ ID NO:4 or SEQ ID NO:6; (b) a nucleotide sequence which hybridizes under conditions of moderate to high stringency to the complement of a second nucleic acid molecule which encodes an amino acid sequence as set forth in SEQ ID NO:2, SEQ ID NO:4 or SEQ ID NO:6; and, (c) a nucleotide sequence which hybridizes under conditions of moderate stringency to the complement of a second nucleic acid molecule as set forth in SEQ ID NO:1, SEQ ID NO:3 or SEQ ID NO:5; and, wherein said nucleic acid molecule encodes an amino acid sequence that has at least about 80% identity to at least one of the amino acid sequences as set forth in SEQ ID NO:2, SEQ ID NO:4 or SEQ ID NO:6.

The present invention also relates to recombinant vectors and recombinant host cells, both prokaryotic and eukaryotic, as well as both stably and transiently transformed/transfected cells, which contain the substantially purified nucleic acid molecules disclosed throughout this specification.

The present invention further relates in part to a substantially purified form of a coccidian CKI protein, as exemplified herein with the identification and disclosure of novel CKI proteins of the *Eimeria* and *Toxoplasma* genera.

The present invention also relates to a substantially purified form of a coccidian CKI protein of the *Eimeria* genus, including but not limited to an *Eimeria* CKI protein from the species *Eimeria tenella*.

The present invention further relates to a substantially purified form of a coccidian CKI protein of the *Toxoplasma* genus, including but not limited to a *Toxoplasma* CKI protein from the species *Toxoplasma gondii*.

The present invention relates to a substantially purified form of a coccidian CKI protein comprising an amino acid sequence selected from group consisting of SEQ ID NO:2, SEQ ID NO:4 and SEQ ID NO:6. Characterization of one or more of these proteins allows for screening methods to identify novel casein kinase I modulators that may have therapeutic activity for animal and/or human health. Thus, the coccidian proteins described herein represent

potential anti-parasitic chemotherapeutic targets for the identification and development of anti-parasitic compounds for the treatment of Toxoplasmosis and coccidiosis.

The present invention further relates to a substantially purified form of a coccidian CKI protein consisting of an amino acid sequence selected from the group consisting of SEQ ID NO:2, SEQ ID NO:4 and SEQ ID NO:6.

A preferred aspect of this portion of the present invention is disclosed in Figure 4 as SEQ ID NO:2, designated EtCKI α , and fully processed forms thereof, which represents a novel coccidian CKI protein from the species *Eimeria tenella*.

Another preferred aspect of this portion of the present invention is disclosed in Figure 4 as SEQ ID NO:4, designated TgCKI α , and fully processed forms thereof, which represents a novel coccidian CKI protein from the species *Toxoplasma gondii*.

Another preferred aspect of this portion of the present invention is disclosed in Figure 4 as SEQ ID NO:6, designated TgCKI β , and fully processed forms thereof, which represents a novel coccidian CKI protein from the species *Toxoplasma gondii*.

The present invention further relates to a substantially purified coccidian CKI protein, said protein comprising at least about 80% amino acid sequence identity with an amino acid sequence selected from the group consisting of SEQ ID NO:2, SEQ ID NO:4 and SEQ ID NO:6.

A preferred aspect of the present invention relates to a substantially purified, fully processed (including any proteolytic processing, glycosylation and/or phosphorylation) and mature coccidian CKI protein obtained from a recombinant host cell (both prokaryotic and eukaryotic, as well as both stably and transiently transformed or transfected) containing a DNA expression vector comprising a nucleotide sequence as set forth in SEQ ID NOs:1, 3 and/or 5, and expresses the *Eimeria* or *Toxoplasma* CKI precursor or mature form of the respective protein. It is especially preferred that the recombinant host cells be a eukaryotic host cell, including but not limited to a mammalian or insect cell line. It is additionally preferable that said coccidian CKI proteins of the present invention be expressed in an inducible eukaryotic expression system.

The present invention also relates to biologically active fragments and/or mutants of the coccidian CKI proteins as initially set forth as SEQ ID NO:2, SEQ ID NO:4 or SEQ ID NO:6, including but not necessarily limited to amino acid substitutions, deletions, additions, amino terminal truncations and carboxy-terminal truncations such that these mutations provide for proteins or protein fragments of diagnostic, therapeutic or prophylactic use and would be useful for screening for agonists and/or antagonists for CKI function.

The present invention further relates to a substantially purified membrane preparation, a partially purified membrane preparations or a cell lysate which has been obtained from a recombinant host cell (both prokaryotic and eukaryotic) transformed or transfected (both stably and transiently) with a DNA expression vector which comprises the nucleic acid molecules of the present invention. These membrane fractions and/or cell lysates will comprise either wild-type or mutant forms of the coccidian CKI proteins of the present invention.

A preferred aspect of the present invention relates to a substantially purified membrane preparation, a partially purified membrane preparations or a cell lysate which has been obtained from a recombinant host cell transformed or transfected with a DNA expression vector which comprises a complete open reading frame as set forth in SEQ ID NOs: 1, 3 and/or 5, expressing a functional form of the respective coccidian CKI protein, either of the *Eimeria* or *Toxoplasma* genera. The subcellular membrane fractions and/or cell lysates from the recombinant host cells (both prokaryotic and eukaryotic, as well as both stably and transiently transformed/transfected cells) contain the functional and processed proteins encoded by the nucleic acids of the present invention. The recombinant-based membrane preparation may comprise a coccidian CKI protein that is essentially free from contaminating proteins, including but not limited to other coccidian source proteins. A preferred aspect of the invention is a membrane preparation or cell lysate which contains one or more coccidian casein kinase I protein(s) comprising the functional form of the CKI proteins as disclosed in Figure 4 (SEQ ID NO:2, designated EtCKI α ; SEQ ID NO:4, designated TgCKI α ; and/or SEQ ID NO:6, designated TgCKI β). These subcellular membrane fractions and/or cell lysates will comprise either wild-type or mutant variations which are biologically functional forms of the *Eimeria* or *Toxoplasma* CKI proteins disclosed herein. It is contemplated that any functional single, homomultimer or heteromultimer combination of the coccidian CKI proteins disclosed herein may be expressed at levels substantially above endogenous levels and hence will be useful in various assays described throughout this specification. A preferred eukaryotic host cell of choice to express the CKI proteins of the present invention is a mammalian or insect based cell line.

The present invention also relates to polyclonal and monoclonal antibodies raised in response to a coccidian CKI protein, including but not limited to CKI proteins of the *Eimeria* and *Toxoplasma* genera disclosed herein, or a biologically active fragment thereof.

The present invention also relates to coccidian CKI fusion constructs, including but not limited to fusion constructs which express at least a portion of the CKI protein linked to various markers, including but in no way limited to the FLAG epitope, GFP (Green fluorescent protein), the MYC epitope, GST, and Fc. Any such fusion constructs may be expressed in the cell line of interest and used to screen for modulators of one or more coccidian CKI proteins.

The present invention relates to methods of expressing coccidian CKI proteins and biological equivalents, including but not limited to CKI proteins of the *Eimeria* and *Toxoplasma* genera, as exemplified by the CKI proteins disclosed herein. The present invention further relates to assays employing coccidian CKI these gene products, as well as recombinant
5 host cells which comprise DNA constructs which express the corresponding proteins. The present invention also relates to compounds identified through the use of coccidian casein kinase I genes and expressed CKI proteins, including agonists or antagonists of coccidian CKI proteins, including but not limited to CKI proteins of the *Eimeria* and *Toxoplasma* genera (e.g., *E. tenella* and *T. gondii*), which act to modulate casein kinase I activity.

10 As used herein, "substantially free from other nucleic acids" means at least 90%, preferably 95%, more preferably 99%, and even more preferably 99.9%, free of other nucleic acids. As used interchangeably with the terms "substantially free from other nucleic acids" or "substantially purified" or "isolated nucleic acid" or "purified nucleic acid," this term also refer to
15 a DNA molecule which comprises a coding region for a coccidian CKI protein, including but not limited CKI proteins of the *Eimeria* or *Toxoplasma* genera, that has been purified away from other cellular components. Thus, a coccidian CKI DNA preparation that is substantially free from other nucleic acids will contain, as a percent of its total nucleic acid, no more than 10%, preferably no more than 5%, more preferably no more than 1%, and even more preferably no more than 0.1%, of non-coccidian CKI nucleic acids. Whether a given coccidian CKI DNA
20 preparation is substantially free from other nucleic acids can be determined by such conventional techniques of assessing nucleic acid purity as, e.g., agarose gel electrophoresis combined with appropriate staining methods, e.g., ethidium bromide staining, or by sequencing.

As used herein, "substantially free from other proteins" or "substantially purified" means at least 90%, preferably 95%, more preferably 99%, and even more preferably 99.9%, free
25 of other proteins. Thus, a coccidian CKI protein preparation, including but not limited to protein preparations of *Eimeria* or *Toxoplasma* CKI proteins, that is substantially free from other proteins will contain, as a percent of its total protein, no more than 10%, preferably no more than 5%, more preferably no more than 1%, and even more preferably no more than 0.1%, of non-coccidian CKI proteins. Whether a given coccidian CKI protein preparation is substantially free
30 from other proteins can be determined by such conventional techniques of assessing protein purity as, e.g., sodium dodecyl sulfate polyacrylamide gel electrophoresis ("SDS-PAGE") combined with appropriate detection methods, e.g., silver staining or immunoblotting. As used interchangeably with the terms "substantially free from other proteins" or "substantially purified," the terms "isolated coccidian CKI protein" or "purified coccidian CKI protein" or "isolated
35 *Eimeria* CKI protein" or "purified *Eimeria* CKI proteins" or "isolated *Toxoplasma* CKI protein"

or "purified *Toxoplasma* CKI protein" also refer to coccidian CKI proteins, including but limited to CKI proteins from the *Eimeria* or *Toxoplasma* genera, that have been isolated from a natural source. Use of the term "isolated" or "purified" indicates that the CKI proteins of the present invention have been removed from their normal cellular environment. Thus, an isolated coccidian CKI protein may be in a cell-free solution or placed in a different cellular environment from that in which it occurs naturally. The term "isolated" does not imply that an isolated coccidian CKI protein is the only protein present, but instead means that a coccidian CKI protein is substantially free of other proteins and non-amino acid material (*e.g.*, nucleic acids, lipids, carbohydrates) naturally associated with the coccidian CKI protein *in vivo*. Thus, for example, a coccidian CKI protein that is recombinantly expressed in a prokaryotic or eukaryotic cell and substantially purified from this host cell which does not naturally (*i.e.*, without intervention) express this CKI protein is, of course, an "isolated coccidian CKI protein" under any circumstances referred to herein. As noted above, a coccidian CKI protein preparation, including but not limited to a protein preparation of *Eimeria* or *Toxoplasma* CKI proteins, that is an isolated or purified coccidian CKI protein will be substantially free from other proteins and will contain, as a percent of its total protein, no more than 10%, preferably no more than 5%, more preferably no more than 1%, and even more preferably no more than 0.1%, of non-coccidian CKI proteins.

As used interchangeably herein, "functional equivalent" or "biologically active equivalent" means a protein which does not have exactly the same amino acid sequence as the corresponding, naturally occurring, coccidian CKI due to alternative splicing, deletions, mutations, substitutions, or additions, but retains substantially the same biological activity as the corresponding, naturally occurring, coccidian CKI. Such functional equivalents will have significant amino acid sequence identity with the naturally occurring coccidian CKI proteins, genes and cDNAs encoding such functional equivalents and can be detected by reduced stringency hybridization with DNA sequences encoding naturally occurring coccidian CKI proteins. For example, a naturally occurring coccidian CKI protein from the *E. tenella* species disclosed herein comprises the amino acid sequence shown as SEQ ID NO:2 and is encoded by SEQ ID NO:1.

As used herein, "a conservative amino acid substitution" refers to the replacement of one amino acid residue by another, chemically similar, amino acid residue. Examples of such conservative substitutions are: substitution of one hydrophobic residue (isoleucine, leucine, valine, or methionine) for another; substitution of one polar residue for another polar residue of the same charge (*e.g.*, arginine for lysine; glutamic acid for aspartic acid).

As used herein, the term "coccidian" refers to a subgroup of the Apicomplexa in which the sexual stage occurs in the gut of the definitive host animal. Infectious diploid oocysts are generated in the gut epithelium and are subsequently released into the environment in the feces. Thus all of the coccidia have an enteric lifecycle stage. *Toxoplasma* is unusual in that the haploid (tachyzoite) stage is not limited to the gut, but can invade and proliferate in any vertebrate cell. As with other coccidia, the sexual stage of *Toxoplasma* occurs in the gut, in this case that of the cat. Examples of other coccidia of clinical or veterinary importance include *Cryptosporidium parvum* (diarrhea in livestock and humans), *Sarcocystis neurona* (neurological symptoms in horses) and *Neospora caninum* (abortions in cattle and neurological symptoms in dogs).

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 shows the nucleotide sequence which encodes a coccidian CKI protein from the species *Eimeria tenella*, as set forth in SEQ ID NO:1, and designated EtCKI α .

Figure 2 shows the nucleotide sequence which encodes a coccidian CKI protein from the species *Toxoplasma gondii*, as set forth in SEQ ID NO:3, and designated TgCKI α .

Figure 3 shows the nucleotide sequence which encodes a coccidian CKI protein from the species *Toxoplasma gondii*, as set forth in SEQ ID NO:5, and designated TgCKI β .

Figure 4 shows a ClustalW alignment of amino acid sequences of parasite CKI proteins, comparing the predicted protein sequences of EtCKI α (SEQ ID NO:2), TgCKI α (SEQ ID NO:4) and TgCKI β (SEQ ID NO:6) with sequences of *Plasmodium falciparum* (PfCKI α ; GenBank Accession No. AF017139; SEQ ID NO:21), *Leishmania major* (LmCKI-2; GenBank Accession No. AAF35365; SEQ ID NO:22) and *Trypanosoma cruzi* (TcCKI-2; GenBank Accession No. AAF00025; SEQ ID NO:23). Peptide fragments identified by microsequencing of affinity purified proteins (Knockaert et al., 2000, *supra*) are highlighted in shaded boxes. TgCKI peptide epitopes, against which antisera were raised, are shown in labeled boxes. N- and C-terminal epitopes are isoform specific (CKI β -Nt, CKI α -Ct, CKI β -Ct). An internal epitope (CKI α -It) was chosen to raise antisera that should recognize all of the aligned CKI isoforms. Conserved amino acids are indicated with a small asterisk below the individual amino acid letter. Brackets are included to delineate the boundaries of the catalytic core region of the enzymes.

Figure 5 shows percentage amino acid identities amongst the sequences shown in Figure 4.

Figure 6 show the transient expression of recombinant coccidian CKI proteins in tachyzoites. Tachyzoites were transfected with 100 μ g of expression plasmid FLAG-TgCKI α (Tg- α), FLAG-TgCKI β (Tg- β), FLAG-EtCKI α (Et- α) or vector alone ('Mock') in triplicate, and

inoculated into cultures of HFF cell monolayers (slides or T-25 flasks). At 24 hours post-infection, anti-FLAG immunofluorescence analyses ("IFA") shows that TgCKI α and EtCKI α are expressed in the cytosol, while TgCKI β exhibits membrane-associated staining (A). At 48 hours post-infection, recombinant enzyme was immunoprecipitated with FLAG agarose from lysates prepared from infected monolayers. Immunoprecipitated enzyme was assayed for kinase activity with casein (B), and detected by Western blot (C) using antisera to internal epitope shared by the different isoforms ('CKI α -It', Figure 4).

Figure 7 shows the mapping of the determinant conferring the membrane association of TgCKI β . TgCKI α , TgCKI β and TgCKI β -CAT fusion constructs are shown schematically with a qualitative assessment of subcellular location ('C' for cytosolic; 'M' for membrane) and casein kinase ('CK') activity indicated. Parasites were transiently transfected with plasmids and the IFA and CK activity determined as described in Figure 6. TgCKI β -CAT fusion proteins were immunoprecipitated with antisera to the CAT marker (shown in black). Blocks of localized sequence similarity between the α and β isoforms of TgCKI (shaded rectangles) were defined using program *AlignX Blocks*, a component of Vector-NTI suite (Informax). Deleted portions of the C-terminal domain of TgCKI β are shown as a thin line.

Figure 8 shows the SDS-PAGE analysis of recombinant TgCKI α and TgCKI β . Enzymes were purified from *E. coli* lysates by FLAG-agarose- and MonoQ ion-exchange chromatography. (A) Lanes 1 and 2 correspond to fractions of peak CK activity from a NaCl gradient and are stained with silver. Bands corresponding to TgCKI α (lane 1, ' α ') and TgCKI β (lane 2, ' β ') are indicated with arrows. Other bands in lane 1 are contaminating *E. coli* proteins. Minor bands in lane 2 are FLAG-TgCKI β degradation products. (B) Western blotting of affinity purified TgCKI α (lane 1) and TgCKI β (lane 2) with antisera to peptide epitopes shown in Figure 4.

Figure 9 shows the partial purification of native TgCKI α from *T. gondii* tachyzoites. Soluble protein was prepared from 2×10^{10} parasites and fractionated by chromatography on a 5 mL HiTrapQ column. CKI phosphopeptide kinase activity was determined in the presence of 1 μ M purvalanol B (triangles) and 200 nM hymenialdisine (open diamonds) or without (closed diamonds) (A). (B) A peak of compound-sensitive CK activity (fraction 7, asterisk) is associated with a 38 kDa band detected with isoform non-selective CK antisera ('CKI α -It'). A second ~60 kDa band is visible in lanes 10-18 that does not correlate with CK activity or cross-react with antisera selective for TgCKI β in duplicate blots (not shown). Kinase activities in fractions 14 and 15 correlate with the presence of CDPK1, which is partially sensitive to purvalanol B and hymenialdisine (see also Table 2). The CKI phosphopeptide is not an optimal substrate for TgCDPK1, so the activity represented underestimates its presence.

Fraction 7 was applied to a 0.1 mL HIC column (C) and 0.1 mL fractions with kinase activity probed with TgCKI α -specific antisera ('CKI α -Ct') (D). Control lanes include column input (MonoQ fraction 7, 'input') and affinity purified recombinant TgCKI α ('FLAGTgCKI α '). Properties of CKI activity in pooled peak fractions (14 and 15) are shown in Tables 1 and 2 ('native CKI α ').

Figure 10 shows the substrate selectivity of recombinant and native TgCKI α . Substrate titrations were performed and $K_{m(app)}$ and $V_{max(app)}$ values determined (see Table 1). For native (B) and recombinant TgCKI α (A), significant levels of concentration-dependent kinase activity were detected only with the synthetic CKI phosphopeptide and bovine casein substrates. The phosphopeptide, KRRRALS(p)VASLPGL (SEQ ID NO:19), yielded two to three fold higher levels of activity than α - or β -casein. Recombinant TgCKI β showed similar properties (Table 1); however, an unphosphorylated CKI peptide, RRKDLHDDEEDEAMSITA (SEQ ID NO:13), also showed some low level activity (C).

Figure 11 shows the partial purification of native EtCKI α from *E. tenella* tachyzoites. Figure 11A shows a silver stained SDS-PAGE gel of fractions (F1-F24) eluted from a MonoS chromatography column, the third of three steps used in the partial purification of the native EtCKI α enzyme. A single ~40 kDa band correlates with peak phosphopeptide kinase activity (B). This kinase activity is sensitive to Compound 20, a (pyrimidyl)(phenyl)substituted fused heteroaryl PKG kinase inhibitor disclosed in PCT International Application PCT/US02/19507 (International publication number WO 03/000682)..

Figure 12 shows preparative 10-20% SDS-PAGE gels stained with Coomassie and Silver for purification of EtCKI α . Bands from fractions 18-20 corresponding to the ~40kDa protein were excised and subjected to tandem LC-MS/MS analysis following digestion with trypsin.

Figure 13 shows results of the tandem LC-MS/MS analysis of the ~40kDa protein isolated from SDS-PAGE gels (see Figure 12). The sequence represents the predicted 39kDa open reading frame of EtCKI α (SEQ ID NO:2). Seven peptide fragments were identified that matched the amino acid sequence of the cloned EtCKI α gene (see shaded boxes). The identity of five of these peptides was confirmed in subsequent methyl ester derivitization experiments.

Figure 14 shows the structure of active purvalanol compounds. Purvalanol B and amino purvalanol share polar phenyl ring substituents (R_1 and R_2) and show good activity against TgCKI α and *T. gondii* grown in vitro, as well as EtCKI α (Table 2). Purvalanol A and structurally related compounds such as Roscovitine and Olomoucine lack these moieties.

Figure 15 shows the structure of the imidazopyridine compound labeled Compound 39 (see Table 3), 4[2-(4-Fluorophenyl)-1-7-(1-methylpiperidin-4-yl)imidazo[1,2-*a*]pyridin-3-yl]pyrimidin-2-amine.

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DETAILED DESCRIPTION OF THE INVENTION

The present invention relates to isolated or purified nucleic acids (polynucleotides) and protein forms which represent coccidian casein kinase I proteins, members of the family of multipotential serine/threonine protein kinases common to all eukaryotic cells and generally referred to herein as the CKI family. This specification discloses three DNA molecules encoding coccidian CKI proteins of the *Eimeria* or *Toxoplasma* genera, specifically CKI proteins from the species *Eimeria tenella* and *Toxoplasma gondii*. The isolated or purified nucleic acid molecules of the present invention, and the proteins they encode, are substantially free from other nucleic acids or proteins, respectively. For most cloning purposes, DNA is a preferred nucleic acid.

The present invention further relates to an isolated or purified nucleic acid molecule (polynucleotide) sequence which encodes a coccidian CKI protein of the *Eimeria* or *Toxoplasma* genera, including but not limited to CKI proteins isolated from the species *Eimeria tenella* and *Toxoplasma gondii*. The *Eimeria* and *Toxoplasma* genera are contained within the phylum *Apicomplexa* which comprises many protozoa of medical and/or veterinary importance. Protozoan species contained within the *Eimeria* and *Toxoplasma* genera generate coccidial infections, labeled coccidiosis in the case of *Eimeria* infections and Toxoplasmosis in the case of *Toxoplasma* infections. Coccidiosis is induced by the intake of a large number of sporulated oocysts by susceptible birds, causing relatively sudden loss of body weight and the onset of clinical signs. While different host organisms are infected by different species of *Eimeria*, poultry are the major hosts for *Eimeria spp.* The United States Department of Agriculture currently recognizes six (6) categories of poultry: chicken, duck, goose, guinea, pigeon and turkey. These different poultry hosts are infected by different species of *Eimeria*, domestic chicken being infected by the following seven species: *E. acervulina*, *E. brunette*, *E. maxima*, *E. mitis*, *E. necatrix*, *E. praecox* and *E. tenella*. Pathogenicities of *Eimeria* infection in chickens range from moderate to severe, *E. tenella* and *E. necatrix* being the most pathogenic. Therefore, one aspect of the present invention relates to purified nucleic acids which encode *Eimeria* CKI proteins isolated from species of *Eimeria* that infect poultry, including but not limited to *E. tenella* and *E. necatrix*. In contrast, the *Toxoplasma* genus is a currently considered a monospecific genus, comprised solely of the species *Toxoplasma gondii*. *T. gondii* is an obligate

intracellular protozoan parasite that infects a broad range of warm-blood animals. In humans, Toxoplasmosis is usually benign; however, in immunocompromised individuals, including cancer and transplant patients and individuals infected with HIV, *T. gondii* infection can result in life-threatening encephalitis, culminating in focal brain disease. Therefore, another aspect of the present invention relates to purified nucleic acids with encode *Toxoplasma* CKI proteins isolated from the species *Toxoplasma gondii*.

The present invention further relates to an isolated or purified nucleic acid molecule (polynucleotide) which encodes a coccidian CKI protein, said nucleic acid molecule comprising or consisting of a nucleotide sequence encoding an amino acid sequence selected from the group consisting of SEQ ID NO:2, SEQ ID NO:4 and SEQ ID NO:6.

One embodiment of the present invention relates to an isolated nucleic acid molecule (polynucleotide) comprising or consisting of a nucleotide sequence encoding the amino acid sequence as disclosed in Figure 4 as SEQ ID NO:2. The amino acid sequence set forth in SEQ ID NO:2 represents the alpha (' α ') isoform of a novel coccidian CKI identified from *Eimeria tenella*, designated EtCKI α . Another embodiment of the present invention relates to an isolated nucleic acid molecule (polynucleotide) comprising or consisting of a nucleotide sequence encoding the amino acid sequences set forth in SEQ ID NO:4 or SEQ ID NO:6. The amino acid sequences set forth in SEQ ID NO:4 and SEQ ID NO:6 represent the alpha (' α ') and beta (' β ') isoforms of novel coccidian CKI identified from *Toxoplasma gondii*, designated TgCKI α and TgCKI β , respectively. The present invention includes codon redundancy which may result in different DNA molecules expressing identical proteins to those disclosed in the present invention.

The present invention also relates to an isolated nucleic acid molecule (polynucleotide) comprising or consisting of a DNA molecule selected from the group consisting of SEQ ID NO:1, SEQ ID NO:3 and SEQ ID NO:5, which encode novel coccidian CKI proteins as set forth in SEQ ID NO:2, SEQ ID NO:4 and SEQ ID NO:6, respectively.

One embodiment of the present invention relates to an isolated or purified nucleic acid molecule (polynucleotide) which encodes mRNA which expresses a novel *Eimeria* CKI protein, this DNA molecule comprising or consisting of the nucleotide sequence disclosed herein as Figure 1 and as set forth in SEQ ID NO:1. The nucleotide sequence set forth in SEQ ID NO:1 encodes the alpha (' α ') isoform of a novel coccidian CKI identified from *Eimeria tenella*, designated EtCKI α , as set forth in SEQ ID NO:2. Another embodiment of the present invention relates to an isolated or purified nucleic acid molecule (polynucleotide) which encodes mRNA which expresses a novel *Toxoplasma* CKI protein, this DNA molecule comprising or consisting of the nucleotide sequence disclosed herein as Figure 2 and as set forth in SEQ ID NO:3. The

nucleotide sequence set forth in SEQ ID NO:3 encodes the alpha (' α ') isoform of a novel coccidian CKI identified from *Toxoplasma gondii*, designated TgCKI α , as set forth in SEQ ID NO:4. A still further embodiment of the present invention relates to an isolated or purified nucleic acid molecule (polynucleotide) which encodes mRNA which expresses a novel
5 *Toxoplasma* CKI protein, this DNA molecule comprising or consisting of the nucleotide sequence disclosed herein as Figure 3 and as set forth in SEQ ID NO:5. The nucleotide sequence set forth in SEQ ID NO:5 encodes the beta (' β ') isoform of a novel coccidian CKI identified from *Toxoplasma gondii*, designated TgCKI β , as set forth in SEQ ID NO:6.

The identification of three novel CKI proteins from *Eimeria tenella* and
10 *Toxoplasma gondii*, coccidian parasites of the *Apicomplexa*, is described in detail in Example 1, said proteins designated as EtCKI α , TgCKI α and TgCKI β . These enzymes have been further characterized as outlined in the Examples Section. These cDNA molecules, as discussed herein, are especially useful as therapeutic targets for the identification and development of anti-parasitic compounds, including but not limited to small molecule agonists and antagonists, for the
15 treatment of Toxoplasmosis and coccidiosis. The cDNAs of the present invention, or portions thereof, are also useful for diagnostic purposes, including but not limited to protein antibody based assays and sequence based assays. In diagnostic assays, the cDNAs would be used to detect infection with said corresponding parasites and the disease states they induce.

The CKI proteins of the present invention are members of the casein kinase I
20 family of serine/threonine protein kinases, a group of highly related, ubiquitously expressed monomeric kinases found in all eukaryotic organisms from protozoa to man. CKI was originally identified for its ability to phosphorylate the milk protein casein; however, this historical name is considered somewhat of a misnomer since milk casein is neither a natural substrate for this family of enzymes nor a particularly optimal one. Seven isoforms have been characterized in
25 mammals, with additional specific homologues identified in lower organisms such as *Saccharomyces cerevisiae*, *Schizosaccharomyces pombe*, *Caenorhabditis elegans*, *Xenopus laevis* and *Drosophila melanogaster*. Among other protein kinases, casein kinase I appears to be unique, having no closely related kinases with similar sequences; however, members of the CKI family are highly related. CKI protein kinases have similar sequence domains consisting of a
30 central kinase domain that is flanked by divergent amino- and carboxyl-terminal regions of variable lengths. CKI proteins preferably phosphorylate substrates having a negatively charged region located upstream of the phosphorylation site, particularly substrates with the recognition motifs S/T(P)-X-X-S/T or D_n-X-X-S/T, wherein S/T(P) represents (phospho)serine or threonine, X is any amino acid, and D is aspartic acid. Importantly, since this recognition motif is relatively
35 ambiguous, many potential substrates for CKI proteins have been identified. Thus, CKI enzymes

have been found to play important and diverse roles within the cell, including contributing to vesicular trafficking, DNA repair, cell cycle progression and cytokinesis in organisms from yeast to humans (Gross, S.D. et al., 1998, *Cell Signal.* 10:699-711; Yu, S. et al., 2002, *Mol. Biol. Cell* 13:2559-2570; Robinson, L.C. et al., 1999, *Mol. Biol. Cell* 10:1077-1092; Behrend, L. et al., 2000, *Eur. J. Cell Biol.* 79:240-251; Vielhaber, E. and Virshup, D.M., 2001, *IUBMB Life.* 51:73-78). CKI enzymes have recently been described for a number of eukaryotic protozoan parasites, including *Plasmodium*, *Leishmania* and *Trypanosoma*. Using affinity chromatography with an immobilized CDK inhibitor (purvalanol B) column, Knockaert et al., 2000, *supra*, identified a 38 kDa protein from *Toxoplasma gondii* extracts with a strong affinity for the purvalanol B matrix. Microsequencing identified the protein as a CKI isoform. The present invention relates in part to novel coccidian CKI proteins from both *T. gondii* (TgCKI α and TgCKI β) and *E. tenella* (EtCKI α).

The present invention also relates to either biologically active fragments or mutants of SEQ ID NO:1 which encode mRNA expressing a novel *Eimeria tenella* CKI protein, designated EtCKI α . Any such biologically active fragment and/or mutant will encode a protein or protein fragment which at least substantially mimics the enzymatic properties of the *E. tenella* CKI protein as set forth in SEQ ID NO:2. Any such polynucleotide includes but is not necessarily limited to nucleotide substitutions, deletions, additions, amino-terminal truncations and carboxy-terminal truncations such that these mutations encode mRNA which express a protein or protein fragment of diagnostic, therapeutic or prophylactic use and is useful for the identification of modulators of CKI activity.

The present invention further relates to either biologically active fragments or mutants of SEQ ID NO:3 which encode mRNA expressing a novel *Toxoplasma gondii* CKI protein, designated TgCKI α . Any such biologically active fragment and/or mutant will encode a protein or protein fragment which at least substantially mimics the enzymatic properties of the *T. gondii* CKI protein as set forth in SEQ ID NO:4. Any such polynucleotide includes but is not necessarily limited to nucleotide substitutions, deletions, additions, amino-terminal truncations and carboxy-terminal truncations such that these mutations encode mRNA which express a protein or protein fragment of diagnostic, therapeutic or prophylactic use and is useful for the identification of modulators of CKI activity.

The present invention still further relates to either biologically active fragments or mutants of SEQ ID NO:5 which encode mRNA expressing a novel *Toxoplasma gondii* CKI protein, designated TgCKI β . Any such biologically active fragment and/or mutant will encode a protein or protein fragment which at least substantially mimics the enzymatic properties of the *T. gondii* CKI protein as set forth in SEQ ID NO:6. Any such polynucleotide includes but is not

necessarily limited to nucleotide substitutions, deletions, additions, amino-terminal truncations and carboxy-terminal truncations such that these mutations encode mRNA which express a protein or protein fragment of diagnostic, therapeutic or prophylactic use and is useful for the identification of modulators of CKI activity.

5 The present invention also relates to isolated or purified nucleic acid molecules which are fusion constructions useful in assays to identify compounds which modulate wild-type coccidian CKI activity, including but not limited to the activity of casein kinase I proteins isolated from the *Eimeria* or *Toxoplasma* genera. Such assays can be used to evaluate the safety and efficacy of specific inhibitors of CKI in host organisms susceptible to coccidial infections. 10 These inhibitors will be useful to treat diseases including coccidiosis and *Toxoplasma* in a range of host organisms, including but not limited to poultry (e.g., domestic chickens) and humans. A preferred aspect of this portion of the invention includes but is not limited to FLAG epitope-tagged CKI fusion constructs. These fusion constructs comprise the open reading frame of the coccidian CKI protein as an in-frame fusion at the NH₂-terminus of the nucleotide sequence encoding the FLAG peptide. Exemplified FLAG epitope-tagged *E. tenella* and *T. gondii* CKI 15 fusion proteins, designated as FLAG-EtCKI α , FLAG-TgCKI α , FLAG-TgCKI β , are described in Example 2 herein. Soluble recombinant FLAG-CKI fusion proteins may be expressed in various expression systems, including *Spodoptera frugiperda* (Sf21) insect cells (Invitrogen) using a baculovirus expression vector (pAcG2T, Pharmingen).

20 The isolated nucleic acid molecule of the present invention may include a deoxyribonucleic acid molecule (DNA), such as genomic DNA and complementary DNA (cDNA), which may be single (coding or noncoding strand) or double stranded, as well as synthetic DNA, such as a synthesized, single stranded polynucleotide. The isolated nucleic acid molecule of the present invention may also include a ribonucleic acid molecule (RNA).

25 The degeneracy of the genetic code is such that, for all but two amino acids, more than a single codon encodes a particular amino acid. This allows for the construction of synthetic DNA that encodes any of the coccidian CKI proteins disclosed herein where the nucleotide sequence of the synthetic DNA differs significantly from the nucleotide sequence of SEQ ID NO:1, SEQ ID NO:3 or SEQ ID NO:5, but still encodes the same coccidian CKI proteins as set 30 forth in SEQ ID NO:2, SEQ ID NO:4 and SEQ ID NO:6, respectively. Such synthetic DNAs are intended to be within the scope of the present invention. If it is desired to express such synthetic DNAs in a particular host cell or organism, the codon usage of such synthetic DNAs can be adjusted to reflect the codon usage of that particular host, thus leading to higher levels of expression of the coccidian CKI protein in the host. In other words, this redundancy in the 35 various codons which code for specific amino acids is within the scope of the present invention.

Therefore, the present invention discloses codon redundancy which may result in differing DNA molecules expressing an identical protein.

It is known that DNA sequences coding for a peptide may be altered so as to code for a peptide having properties that are different than those of the naturally occurring peptide.

5 Methods of altering the DNA sequences include but are not limited to site directed mutagenesis. Examples of altered properties include but are not limited to changes in the affinity of an enzyme for a substrate or a receptor for a ligand.

As used herein, "purified" and "isolated" are utilized interchangeably to stand for the proposition that the nucleic acid, protein, or respective fragment thereof in question has been
10 substantially removed from its *in vivo* environment so that it may be manipulated by the skilled artisan, such as but not limited to nucleotide sequencing, restriction digestion, site-directed mutagenesis, and subcloning into expression vectors for a nucleic acid fragment, as well as obtaining the protein or protein fragment in pure quantities so as to afford the opportunity to generate polyclonal antibodies, monoclonal antibodies, amino acid sequencing, and peptide
15 digestion. Therefore, the nucleic acids claimed herein may be present in whole cells or in cell lysates or in a partially purified or substantially purified form. A nucleic acid is considered substantially purified when it is purified away from environmental contaminants. Thus, a nucleic acid sequence isolated from cells is considered to be substantially purified when purified from cellular components by standard methods, while a chemically synthesized nucleic acid sequence
20 is considered to be substantially purified when purified from its chemical precursors.

A preferred aspect of the present invention is disclosed in Figure 1 and SEQ ID NO:1, a coccidian cDNA encoding an *Eimeria tenella* casein kinase I alpha gene, EtCKI α , disclosed as follows:

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1  GCGGCCGCGT  CGACGTCTTT  GCTGCCGCAC  AGGGAGCAGC  AGCAGCCGCC  GACCCGATCC
25  61  CTTGGGAGCC  CACCAAGTGC  TGCCTTGCT  TAGCAGCTAC  AGGAGCTGCC  GCGGGGTGCG
    121  TCCCTGAGGC  AGCGTGCATG  TATGGTCCGG  CAGCCAGCTT  GGTGTCGCAG  CCGTACTTCT
    181  TGGAAGCGAG  AGAGACTGTG  GGAGAGCGCA  AATCACTCCA  GCCGCTTCCA  GGGGAGTCTG
    241  GGGACCGCAG  GAGCGTTGGA  GGCTGCCTGC  CGGCATAAAC  AGGAACAAGC  GCATTCTTAT
    301  TCTTCTGTGG  TTGCTGAGTT  CTGGCTGCGT  TCAAGGGGGT  TCACCTCTTC  CCCTTCTGGC
30  361  GAGTTTTTGC  TCGCTCTTTC  CCTAAGAAGC  AGCGCCACGT  GCGTGCGGTG  CCTCAGCCTG
    421  ACGCGGTGCA  CCTTTTACGT  AAGAGCGTCG  ATAGCATCGG  TCATCTACAG  CAGCGTGCTG
    481  CTGCTTCCGT  GACCTTTACA  CTGCTTGTGG  CGGGCCGTCT  TGTAGAGGGG  CCATCTGCTT
    541  GTTCGCTGCT  GGACGCAGAC  CCGGCGCCCG  ACATTTCGGG  CAGCCGGGCA  GTTGAGATAA
    601  ACCGGCTGCC  CGGTGGCCGT  CGAAATTGAA  GCAGGATCTC  TACAGTAAGG  AACAAATCGC
35  661  GCTATTTTTA  AGGAGTGTGT  ATACTTGGGG  CGTTACTCGT  GAGTATTGCT  GATGATGGAC

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721 GTCCGTGTGG GGGGTAAGTA TCGTTTGGGG AGGAAGATTG GGAGCGGATC CTTCGGCGAC
 781 ATCTACCTTG GTACGAACAT CTCAACAGGA GATGAAGTCG CTATCAAATT GGAAAGCGTG
 841 CGGTCTAGGC ATCCACAACCT AATCTATGAA AGCAAGCTGT AAAAAATCCT AACGGGTGGA
 901 ATCGGAATCC CGACTCTTTA CTGGTATGGG ATCGAGGGGG ATTACAACGT TATGATTATT
 5 961 GAGCTTTTGG GCCCGTCTCT TGAGGACCTC TTCAGCATTT GCAACAGAAA GCTTTCCTTG
 1021 AAGACTGTTT TGATGCTCGC CGACCAAATG CTAAATCGTA TTGAGTTCGT CCACAGCAGA
 1081 CATTTTCATCC ATCGAGACAT CAAGCCTGAC AATTTTTTGA TCGGTAGGGG CAAAAAGATG
 1141 TCCATTGTTT TTGCTATCGA CTTTGGCCTC GCAAAGAAGT ACAGAGATCC CAGAACACAG
 1201 TCCCATATTC CTTATCGAGA AGGGAAGAAC CTGACAGGTA CCGCGAGGTA CGCCTCTGTG
 10 1261 AACACCCACT TGGGAATAGA ACAGAGCAGG CGCGATGATC TGGAGCGCT CGGCTACGTC
 1321 TTAATGTACT TCAACAGAGG TTCCTTACCC TGGCAAGGAT TAAAGGCCAC TACGAAGAAA
 1381 GATAAATATG ACAAGATTAT GGAGAAGAAG ATGTCCACCC CTATTGAAGT CCTTTGCAAA
 1441 CAATTTCCAT TTGAGTTTAT CACATATCTG AACTATTGCC GGTCTCTGCG ATTCGAAGAT
 1501 CGCCCGGACT ATTCCTATTT GAGACGGTTG TTCAAGGATC TTTTCTTCCG TGAGGGATAC
 15 1561 CAGTATGACT TTATATTCGA TTGGACATTT CTGCATGCTG AGAGAGAGCG CGAGCGTCAA
 1621 AGACGATCGA TGGTCAACCA AGGCGCAGAA TCAGGGAACC AGTGGAGACG AGACGCGTCG
 1681 GGCAGAGATC CACTTGGACG GTTGCCTCAG TTAGAACCGT AATCTCTTTA CGGGCAGATT
 1741 GCCGTACGGG TCTTCTGCTC ATTCAGTGGC AGTGCCACCG CAGTGTCTATC TGAGGCTGTG
 1801 GCTTCAGGAT GTGGTAGCCA GTTACCATGG TCACTTGCCC TCGCTAGGAC AGCCTTCGCA
 20 1861 GGGAAATGTC ACAGTAGCCT GCATTATGTG GTGTGAGAAC TGCTAGCGCA TTCCTGTAGT
 1921 TGCTTTTACG AAGCAGGATA CGCAGCGTGC ATCACGCGGT GGTTCGAGCG CTCGCTACGC
 1981 ATCACAGGGC TGTGAGGCAA GTTAGTATCT TTGGGGGACG AGTTGAGAGT GTCAGAATCG
 2041 ATAGTCTCAG GGCATGCAGG CGAAATGGAG GCTGCGCCAG TAGTGCCAGC CGGTGGCGAA
 2101 GCGCTCAAAAT TTACTTTTTT TGTGCTGGG GATATTGTTA GAGCAACAAC TTGGGTCTAG
 25 2161 ATGCTACTGA TAAAAA AAAA (SEQ ID NO:1).

Another preferred aspect of the present invention is disclosed in Figure 2 and SEQ
 ID NO:3, a coccidian cDNA encoding a *Toxoplasma gondii* casein kinase I alpha gene, TgCKI α ,
 disclosed as follows:

1 CCTCGTTTTG CTTCAATCCC CGCCTTTTCT CTGTAGCTAA CCAAAGGAAC AAAGTCAGCG
 30 61 GTAGAAGCCG TTTCTTCTGT CCGCTTCCCA CTCTTCCCGT TCGGCTGCCC CTGCAGAGCG
 121 CCCTTTCTAT GCGTTGCCAC CCGTCTGCAA GTATCGCGTC TTTGCTCTCA TCAGTGATTT
 181 TCTTTGCGTG TCGCGTTCGG GACGCCCTTT TCTCTCCTCA ACTAACTAGC AGACGTTTCT
 241 TCCGTCCCGC ATGCGACAGC GAAGGGCAGC TCCCCCAGT TCTTCATCGC CCACCTGTTG
 301 TGCAACTTGT CGCCCGTCGT TCTTCACTTC TTCTCTCCCA TCCTCTCGTG ACTCTTCCCTC
 35 361 TCGAGAACTC TTTCTGTCTGA ACTCTCAACC CCCACGACTG CTGGTTTCGT GGCCGTCCCG

421 CATGCACCTT GTGTCCCGCC GCCTTGGCGC AAACACCCGC TTTCTCTGCT GTCCGCCTCC
 481 CGGTGGACTT CTCTCCGTGT TTTTTCGTGT TGCCAAAAGT TTGTCTGCTT TGACGTTTCT
 541 CTGCTCACCC ATTGCCCCGCT CTTGATGAGG AACGCTCCAC ATTGACAGCG AACTCACAGC
 601 ACGCACCCCTC CGCGAGCGGA CTTTCACGAG CGAGGCAAGA ATCCATCGTC ACCCCGCCTA
 5 661 CACGTACACT ACTCCACTTG GGTGCCCACG CGCGGCTTCT GGGAGACAGA GACGGTCCCTC
 721 GTTTTCCGTG TCAGAACTTT GTCGAGGAAA CGCTGCTGCT GGCAGCGGGG ATTGTGACCC
 781 CCCTCGGCGA ACGGGCGAAG CCGCCCTGTC GCGCGTCGCG ACTCAGCTGA GGCAGACGGC
 841 GGTGCGCGGC GTGACCTCTC TTTCTTTTGT CATTCGGCCC TGATTGCAGC ACGAAGGATG
 901 GAGGTCAGGG TCGGAGGCAA GTACCGACTT GGTGGAAGA TCGGCAGCGG GTCATTCCGT
 10 961 GATATTTATA TCGGTGCAAA CATTTTGACG GGGGATGAGG TGGCGATCAA GTTGGAGTCT
 1021 ATCAAGTCGA AGCACCCGCA GCTGCTCTAT GAGTCGAAGC TGTACAAACT GCTGGCTGGC
 1081 GGCATTGGGA TTCCCATGGT CCACTGGTAC GGCATCGAAG GAGACTACAA TGTTATGGTT
 1141 ATCGACCTTC TCGGCCCTTC TCTGGAGGAC CTTTTCAGTA TCTGCAATCG CAAACTCTCT
 1201 CTCAAGACGG TGTTGATGCT CGCAGACCAG ATGCTCAACC GCATCGAGTT TGTCCATAGC
 15 1261 AAGAACTTCA TCCATCGCGA TATCAAACCC GACAACTTCC TCATTGGCCG TGGAAAGAAG
 1321 ATGTCCGTCG TCTACATCAT CGATTTCCGT TTGGCAAAGA AATATCGAGA CCCAAAGACT
 1381 CAGCAACATA TCCCATACAG GGAAGGCAAG AACCTAACAG GCACAGCGCG TTACGCTTCC
 1441 ATCAACACCC ACCTGGGGAT CGAGCAGAGT CGGCAGACG ACCTAGAGGC GCTCGGTTAC
 1501 GTTCTCATGT ACTTCAATAG AGGTTCTCTT CCGTKGCAGG GTCTGAAGGC GACGACGAAG
 20 1561 AAGGACAAAT ACGACAAGAT TATGGAGAAG AAAATGTCTA CTCCCATCGA AATTTTGTGC
 1621 AAGCATTTCC CATTCGAGTT CATCACCTAC TTGAATTACT GCCGGTCCCT GCGCTTCGAG
 1681 GATCGTCCTG ACTACGCATA CTTGCGACGC CTGTTCAAAG ACTTGTTTTT TAGAGAGGGA
 1741 TATCAGTACG ACTTCATCTT CGACTGGACT TTCATCAACA CGGAGAAGGA TCGCGCGAGT
 1801 CGAAGAAGCC AGCAAGTTTA TGTGGAAGAC AACCGGCAAG TTGAGGAGAA TCAGAACGAG
 25 1861 TTGCCGATGT AGGGTGGTCG GTGTGCGGAG GCCGGCGGGG AGCGTGGAGT CCGCTGAGTC
 1921 TGGAAGTCTG CAGACTGTGC TCTGGCACTC GACCCACTTG TTTGTGTTTC CCTCGACTCG
 1981 CGCAGGTCGA GGAACACAGA GACGAACAGG TTACCCAGGA GTGTTTTTGG TCAGGACGCG
 2041 CGTCTCCCTC TGAGTTTCGC AAAGTTGCCC CTGGAA (SEQ ID NO:3).

Another preferred aspect of the present invention is disclosed in Figure 3 and SEQ
 ID NO:5, a coccidian cDNA encoding a *Toxoplasma gondii* casein kinase I beta gene, TgCKI β ,
 disclosed as follows:

1 TTAACCCTCA CTAAAGGGAA CAAAAGCTGG AGCTCCACCG CGGTGGCGGC GCACCGAGGA
 61 AAACGCAGCT CGTAAGAGAC AGTTCTCTCG GTGAGAAGAG CTATCCGAGA AGGACACCAT
 121 GCGCACCAT CAAGACACCC GCAACCACAC GGGGGTCGGA CCTCTTCTGT CTATCCCTCT
 35 181 GAAAGATTTG AAGATCGCCG GCGTCTGGAA AATCGGCAGA AAAATCGGAT CCGGTTCCCT

241 CGGCGACATA TACAAAGGCC TGAATTCTCA GACCGGTCAG GAGGTGGCGC TGAAGGTCTGA
301 AAGCACCAAG GCGAAGCATC CGCAGTTGCT GTACGAATAC AAACTTTTGA AGCATTGCA
361 GGGAGGAACG GGCATTGCTC AAGTGTCTG TTGCGAGACT GCGGGCGACC ATAACATCAT
421 GGCCATGGAG TTGCTCGGAC CTTCTTTAGA GGACGTCTTC AACTTGTGCA ATCGCACCTT
5 481 CTCTCTCAA ACCATTCTTC TTCTCGCCGA CCAGTTTCTG CAACGCGTCG AGTACATCCA
541 CTCCAAGAAT TTCATTCA CA GAGATATCAA ACCAGATAAC TTCTTTCTCG GCGGTGCCGG
601 CAATCAAAAC ACGATCTACG TGATCGACTT CGGCCTGGCG AAGAAGTTTC GCGATCCGAA
661 AACGCACCA CATATTCCGT ACAGAGAAAA CAAGAATCTC ACGGGAACGG CGCGCTACGC
721 GTCCATCAGT GCGCATCTGG GTTCCGAGCA GAGTCGCCGA GATGACCTCG AAGCAGTCGG
10 781 CTACGTTCTC ATGTACTTCT GTCGAGGAG CACGCTGCCT TGGCAGGGCA TCAAAGCGAA
841 TACCAACAG GAGAAGTACC ACAAGATCAT GGAGAAGAAG ATGTCGACGC CCGTCGAGGT
901 GCTATGCAAG GGATATCCAA GCGAATTTGC CACATACTTG CACTACTGCC GCTCCTTGCG
961 ATTCGAGGAC CGACCGGACT ACGCCTACCT CAAGCGACTC TTTCGAGATC TCTACATCAA
1021 AGAGGGCTAC GATGACAGTG ACCGCGAATT CGACTGGACA GTGAAACTTT CGTCGCGCAG
15 1081 TCTCGGACCG CCAAGCAGTC GAGCGCAACA TGTTTTACTG AGTCAAGACA CCCGAACGCG
1141 AGGGAAGCGG GAGACAGATC GACCTGTCGC TCGCGGAGT GGCGACCGCG AACGAGGAAT
1201 CCATTTACAG AACGGGAACG TGGGCAATCC TTCGATGGCA ACGAACCCCG GCGGCCTGTC
1261 AGTCATGGTG CATGAACGCA CGAGTCTGGT GGATCAGGGA GACCGTGGGT CGCGCGAAAC
1321 TTCTACGCGG AAAGAAGACG CGAAGGACGG CAGATGGCCA GGAGGCAGAT TTTCTTGCT
20 1381 TCCACTGTTA TGTCGGCGCT CTCCGACGAA GGCCTAGATG AACTGCGGAG GCGCTCCTGT
1441 CCCCAGAGT GGCATCTCTC TCCTTCATTG TCGTTGTTCC CCTGCAACTC GAGTCCACCC
1501 TTGACATCCT CGTCTCTCTC TTCTGTGCG TTTCTCTTT CTCTCTCTCT CCCCCCTAGC
1561 TTCGTTCTCT CTTTCTATC CTGCTTCGGC GTCGCCCTAC TTCTCTCTCT ACTTCTCTCC
1621 CTTTGTGTTT TCTTCGCGGC GTCTCTCCTT CACTCTGTCT CCGCCTCTGA CGCCGCGCGG
25 1681 GAGCCGTTTC CTGCAGGCAG CTCAGGCAAT ACCTGCCTGC AGGTGCCTCT CTTTTTTGAG
1741 CGTCTCTCTT TCCTCGTCGA AACGGTCCTC ACAGCTTCCT CTCCCTGGGG ACGCCGTGGG
1801 CGTAAGTTCT TTTTTTGACG GTCCCGGTGG GCTGGCGTTG TTCGCCTGCC TTCCGCGCAT
1861 GCACTCCGAG CATTTTTGCC TGGCCTGGAC TTCTCCGAGC GAGAGTTGCG GTTTGGCTTC
1921 TGCATCGTCT CCTGCGCTGC TTTCATTCT CTAGGTTTCT GCTTGC GGCC TCCGTGTACA
30 1981 GAAATCGGAA GGTGAAGGCG TAGTGGCCAG AGAACGAAGC AAACGAGAGA ACCACGTTCC
2041 ACCTTGTCG CACGCATGCA TCTACGCATG CACGGTATTT AAGCCGATTT TTTGTGTATG
2101 TATATAGATG TATATATATA TGTATCTACA TGTATCTACC TATATATATG TGTGTGTGTA
2161 AGTGGAAAGT TATTTTTGCA TGTGCAGAAA GCTTTCTTTT CCGCTGGCAT GCTGGAAGAA
2221 GGGCAGGAGG CGACGATCCT GCGAGTCAGG GCGTTCCCTT GTTCCAGTG AGTTAACCGA

2281 ATTGTTTATT GATATGCGTT TGCATGCATC GACAATGGAT CCTAGACACG CCCGTTTAAA
 2341 ATCAGAGGTA TTCCTAAAAA AAAAAAAAAA AAA (SEQ ID NO:5).

The above-exemplified isolated DNA molecules, shown in Figures 1-3, comprise the following characteristics:

5 EtCKI α (SEQ ID NO:1): 2182 nucleotides; initiating Met at nuc. 715-717 and "TAA" termination codon at nuc.1720-1722; open reading frame results in an expressed protein of 335 amino acids, as set forth in SEQ ID NO:2.

10 TgCKI α (SEQ ID NO:3): 2076 nucleotides; initiating Met at nuc. 898-900 and "TAG" termination codon at nuc. 1870-1872; open reading frame results in an expressed protein of 324 amino acids, as set forth in SEQ ID NO:4.

TgCKI β (SEQ ID NO:5): 2373 nucleotides; initiating Met at nuc. 119-121 and "TAG" termination codon at nuc. 1415-1417; open reading frame results in an expressed protein of 432 amino acids, as set forth in SEQ ID NO:6).

15 The percent identity of the amino acid sequences encoded by the exemplified cDNA molecules of the present invention to both each other as well as other eukaryotic protozoan parasite CKI isoforms are as follows:

20 TgCKI α and TgCKI β - 48%;
 TgCKI α and PfCKI α - 68%;
 TgCKI α and LmCKI-2 - 58%;
 TgCKI α and TcCKI-2 - 62%;
 TgCKI β and PfCKI α - 45%;
 TgCKI β and LmCKI-2 - 44%;
 TgCKI β and TcCKI-2 - 42 %;
 EtCKI α and TgCKI α - 81%;
 25 EtCKI α and TgCKI β - 48%;
 EtCKI α and PfCKI α - 67%;
 EtCKI α and LmCKI-2 - 59%; and,
 EtCKI α and TcCKI-2 - 60%.

30 To determine the percent identity of two amino acid sequences or two nucleic acid sequences, the sequences are aligned for optimal comparison purposes (*e.g.*, gaps can be introduced in one or both of a first and a second amino acid or nucleic acid sequence for optimal alignment, and non-homologous sequences can be disregarded for comparison purposes). In one embodiment of the invention, the length of a reference sequence aligned for comparison purposes is at least 30%, 40%, 50%, 60% or 70% of the length of the reference sequence, more preferably
 35 at least 80%, and most preferably at least 90% or more of the length of the reference sequence.

The amino acid residues or nucleotides at corresponding amino acid positions or nucleotide positions are then compared. When a position in the first sequence is occupied by the same amino acid residue or nucleotide as the corresponding position in the second sequence, then the molecules are identical at that position (as used herein, amino acid or nucleic acid "identity" is equivalent to amino acid or nucleic acid "homology"). The percent identity between the two sequences is a function of the number of identical positions shared by the sequences, taking into account the number of gaps, and the length of each gap, which need to be introduced for optimal alignment of the two sequences.

The comparison of sequences and determination of percent identity and similarity between two sequences can be accomplished using a mathematical algorithm (*see e.g.*, *Computational Molecular Biology*, Lesk, A. M., ed., Oxford University Press, New York, 1988; *Biocomputing: Informatics and Genome Projects*, Smith, D. W., ed., Academic Press, New York, 1993; *Computer Analysis of Sequence Data, Part 1*, Griffin, A. M., and Griffin, H. G., eds., Humana Press, New Jersey, 1994; *Sequence Analysis in Molecular Biology*, von Heinje, G., Academic Press, 1987; and *Sequence Analysis Primer*, Gribskov, M. and Devereux, J., eds., M Stockton Press, New York, 1991). In a preferred embodiment, and as exemplified in calculating the percent identities between the coccidian amino acid sequences disclosed herein, the percent identity between two amino acid sequences is determined based on alignments generated with the Clustal W algorithm (Thompson, J.D. et al., 1994, *Nucleic acids Res.* 22:4673-4680). This algorithm is incorporated into many commercial software packages, in this case the alignX software program in the Vector NTI suite (version 8.0). Default Clustal W parameters were used to generate pairwise alignments from which percent identity values were calculated (gap opening penalty of 10; gap extension penalty of 0.1). The percent identity is defined as the number of identical residues divided by the total number of residues and multiplied by 100. If sequences in the alignment are of different lengths (due to gaps or extensions), the length of the longest sequence will be used in the calculation, representing the value for total length.

To this end, the present invention further relates to an isolated or purified nucleic acid molecule encoding a coccidian CKI protein, including but not limited to a CKI protein of the *Eimeria* or *Toxoplasma* genera, wherein the nucleic acid molecule comprises a nucleotide sequence selected from the group consisting of: (a) a nucleotide sequence which encodes an amino acid sequence as set forth in SEQ ID NO:2, SEQ ID NO:4 and/or SEQ ID NO:6; (b) a nucleotide sequence which hybridizes under conditions of moderate to high stringency to the complement of a second nucleic acid molecule which encodes an amino acid sequence as set forth in SEQ ID NO:2, SEQ ID NO:4 and/or SEQ ID NO:6; and, (c) a nucleotide sequence which hybridizes under conditions of moderate stringency to the complement of a second nucleic acid

molecule as set forth in SEQ ID NO:1, SEQ ID NO:3 and/or SEQ ID NO:5; and, wherein said nucleic acid molecule encodes an amino acid sequence that has at least about 80% identity to at least one of the amino acid sequences as set forth in SEQ ID NO:2, SEQ ID NO:4 and/or SEQ ID NO:6.

5 One embodiment of the present invention relates to an isolated or purified nucleic acid molecule encoding a coccidian CKI protein, including but not limited to a CKI protein of the *Eimeria* or *Toxoplasma* genera, wherein the nucleic acid molecule comprises a nucleotide sequence selected from the group consisting of: (a) a nucleotide sequence which encodes an amino acid sequence as set forth in SEQ ID NO:2 and/or SEQ ID NO:4; (b) a nucleotide
10 sequence which hybridizes under conditions of moderate to high stringency to the complement of a second nucleic acid molecule which encodes an amino acid sequence as set forth in SEQ ID NO:2 and/or SEQ ID NO:4; and, (c) a nucleotide sequence which hybridizes under conditions of moderate stringency to the complement of a second nucleic acid molecule as set forth in SEQ ID NO:1 and/or SEQ ID NO:3; and, wherein said nucleic acid molecule encodes an amino acid
15 sequence that has at least about an 80% identity to at least one of the amino acid sequences as set forth in SEQ ID NO:2 and/or SEQ ID NO:4.

Another embodiment of the present invention relates to an isolated or purified nucleic acid molecule encoding a coccidian CKI protein, including but not limited to a CKI protein of the *Eimeria* or *Toxoplasma* genera, wherein the nucleic acid molecule comprises a
20 nucleotide sequence selected from the group consisting of: (a) a nucleotide sequence which encodes an amino acid sequence as set forth in SEQ ID NO:4 and/or SEQ ID NO:6; (b) a nucleotide sequence which hybridizes under conditions of moderate to high stringency to the complement of a second nucleic acid molecule which encodes an amino acid sequence as set forth in SEQ ID NO:4 and/or SEQ ID NO:6; and, (c) a nucleotide sequence which hybridizes
25 under conditions of moderate stringency to the complement of a second nucleic acid molecule as set forth in SEQ ID NO:3 and/or SEQ ID NO:5; and, wherein said nucleic acid molecule encodes an amino acid sequence that has at least about an 80% identity to at least one of the amino acid sequences as set forth in SEQ ID NO:4 and/or SEQ ID NO:6.

A further embodiment of the present invention relates to an isolated or purified
30 nucleic acid molecule encoding a coccidian CKI protein, including but not limited to a CKI protein of the *Eimeria* or *Toxoplasma* genera, wherein the nucleic acid molecule comprises a nucleotide sequence selected from the group consisting of: (a) a nucleotide sequence which encodes an amino acid sequence as set forth in SEQ ID NO:2 and/or SEQ ID NO:6; (b) a nucleotide sequence which hybridizes under conditions of moderate to high stringency to the
35 complement of a second nucleic acid molecule which encodes an amino acid sequence as set

forth in SEQ ID NO:2 and/or SEQ ID NO:6; and, (c) a nucleotide sequence which hybridizes under conditions of moderate stringency to the complement of a second nucleic acid molecule as set forth in SEQ ID NO:1 and/or SEQ ID NO:5; and, wherein said nucleic acid molecule encodes an amino acid sequence that has at least about an 80% identity to at least one of the amino acid sequences as set forth in SEQ ID NO:2 and/or SEQ ID NO:6.

One embodiment of the present invention relates to an isolated or purified nucleic acid molecule encoding a coccidian CKI protein, including but not limited to a CKI protein of the *Eimeria* or *Toxoplasma* genera, wherein the nucleic acid molecule comprises a nucleotide sequence selected from the group consisting of: (a) a nucleotide sequence which encodes an amino acid sequence as set forth in SEQ ID NO:2; (b) a nucleotide sequence which hybridizes under conditions of moderate to high stringency to the complement of a second nucleic acid molecule which encodes an amino acid sequence as set forth in SEQ ID NO:2; and, (c) a nucleotide sequence which hybridizes under conditions of moderate stringency to the complement of a second nucleic acid molecule as set forth in SEQ ID NO:1; and, wherein said nucleic acid molecule encodes an amino acid sequence that has at least about an 80% identity to at least one of the amino acid sequences as set forth in SEQ ID NO:2.

Another embodiment of the present invention relates to an isolated or purified nucleic acid molecule encoding a coccidian CKI protein, including but not limited to a CKI protein of the *Eimeria* or *Toxoplasma* genera, wherein the nucleic acid molecule comprises a nucleotide sequence selected from the group consisting of: (a) a nucleotide sequence which encodes an amino acid sequence as set forth in SEQ ID NO:4; (b) a nucleotide sequence which hybridizes under conditions of moderate to high stringency to the complement of a second nucleic acid molecule which encodes an amino acid sequence as set forth in SEQ ID NO:4; and, (c) a nucleotide sequence which hybridizes under conditions of moderate stringency to the complement of a second nucleic acid molecule as set forth in SEQ ID NO:3; and, wherein said nucleic acid molecule encodes an amino acid sequence that has at least about an 80% identity to at least one of the amino acid sequences as set forth in SEQ ID NO:4.

A further embodiment of the present invention relates to an isolated or purified nucleic acid molecule encoding a coccidian CKI protein, including but not limited to a CKI protein of the *Eimeria* or *Toxoplasma* genera, wherein the nucleic acid molecule comprises a nucleotide sequence selected from the group consisting of: (a) a nucleotide sequence which encodes an amino acid sequence as set forth in SEQ ID NO:6; (b) a nucleotide sequence which hybridizes under conditions of moderate to high stringency to the complement of a second nucleic acid molecule which encodes an amino acid sequence as set forth in SEQ ID NO:6; and, (c) a nucleotide sequence which hybridizes under conditions of moderate stringency to the

complement of a second nucleic acid molecule as set forth in SEQ ID NO:5; and, wherein said nucleic acid molecule encodes an amino acid sequence that has at least about an 80% identity to at least one of the amino acid sequences as set forth in SEQ ID NO:6.

"Stringency" of hybridization reactions is readily determinable by one of ordinary skill in the art, and generally is an empirical calculation dependent upon probe length, washing temperature, and salt concentration. In general, longer probes require higher temperatures for proper annealing, while shorter probes need lower temperatures. Hybridization generally depends on the ability of denatured DNA to re-anneal when complementary strands are present in an environment below their melting temperature. The higher the degree of desired homology between the probe and hybridizable sequence, the higher the relative temperature which can be used. As a result, it follows that higher relative temperatures would tend to make the reaction conditions more stringent, while lower temperatures less so. Generally, stringent conditions are selected to be about 5°C lower than the thermal melting point (T_m) for the specific sequence at a defined ionic strength and pH. The T_m is the temperature (under defined ionic strength, pH and nucleic acid concentration) at which 50% of the probes complementary to the target sequence hybridize to the target sequence at equilibrium. Since target sequences are generally present at excess, at T_m , 50% of the probes are occupied at equilibrium. Typically, stringent conditions will be those in which the salt concentration is less than about 1.0 M sodium ion, typically about 0.01 to 1.0 M sodium ion (or other salts) at pH 7.0 to 8.3; and the temperature is at least about 30°C for short probes, primers or oligonucleotides (e.g., 10 to 50 nucleotides) and at least about 60°C for longer probes, primers and oligonucleotides. Stringent conditions may also be achieved with the addition of destabilizing agents, such as formamide. For additional details and explanation of stringency of hybridization reactions, see Ausubel et al., *Current Protocols in Molecular Biology*, Wiley Interscience Publishers, (1995).

Conditions of "high stringency," as defined herein, may be identified by those that: (1) employ low ionic strength and high temperature for washing, for example 0.015 M sodium chloride/0.0015 M sodium citrate, 0.1% sodium dodecyl sulfate (SDS) at 50°C; (2) employ during hybridization a denaturing agent, such as formamide, for example, 50% (v/v) formamide with 0.1% bovine serum albumin (BSA), 0.1% Ficoll, 0.1% polyvinylpyrrolidone, 50 mM sodium phosphate buffer at pH 6.5 with 750 mM sodium chloride/750 mM sodium citrate at 42°C; or (3) employ 50% formamide, 5X SSC (0.75 M NaCl, 0.075 M sodium citrate), 50 mM sodium phosphate (pH 6.8), 0.1% sodium pyrophosphate, 5X Denhardt's solution, sonicated salmon sperm DNA (50 µg/ml), 0.1% SDS, and 10% dextran sulfate at 42°C, with washes at 42°C in 0.2X SSC (sodium chloride/sodium citrate) and 50% formamide at 55°C, followed by a high-stringency wash consisting of 0.1X SSC containing EDTA at 55°C.

"Moderate stringency" conditions may be identified as described by Sambrook et al., *Molecular Cloning: A Laboratory Manual*, New York: Cold Spring Harbor Press, 1989, and include the use of washing solution and hybridization conditions (e.g., temperature, ionic strength and percent SDS) less stringent than those described above. An example of moderately stringent conditions is overnight incubation at 37°C in a solution comprising: 20% formamide, 5X SSC (150 mM NaCl, 15 mM trisodium citrate), 50 mM sodium phosphate (pH 7.6), 5X Denhardt's solution, 10% dextran sulfate, and 20 mg/ml denatured sheared salmon sperm DNA, followed by washing the filters in 1X SSC at about 37-50°C. The skilled artisan will recognize how to adjust the temperature, ionic strength, etc. as necessary to accommodate factors such as probe length and the like. An example of progressively higher stringency conditions is as follows: 2X SSC/0.1% SDS at about room temperature (hybridization conditions); 0.2X SSC/0.1% SDS at about room temperature (low stringency conditions); 0.2X SSC/0.1% SDS at about 42°C (moderate stringency conditions); and 0.1X SSC at about 68°C (high stringency conditions). Washing can be carried out using only one of these conditions, e.g., high stringency conditions, or each of the conditions can be used, e.g., for 10-15 minutes each, in the order listed above, repeating any or all of the steps listed. However, as mentioned above, optimal conditions will vary, depending on the particular hybridization reaction involved, and can be determined empirically.

The present invention also relates to recombinant vectors and recombinant host, both prokaryotic and eukaryotic, which contain the substantially purified nucleic acid molecules disclosed throughout this specification.

The present invention relates to purified forms of coccidian casein kinase I proteins substantially free from other proteins, including but not limited to coccidian CKI proteins of the *Eimeria* or *Toxoplasma* genera. One aspect of the present invention relates to substantially purified *Eimeria* CKI proteins isolated from species of *Eimeria* that infect poultry, including but not limited to *E. tenella* and *E. necatrix* which are highly pathogenic in domestic chickens. Another aspect of the present invention relates to substantially purified *Toxoplasma* CKI proteins isolated from the species *Toxoplasma gondii*.

The present invention further relates to substantially purified coccidian CKI proteins which comprise or consist of an amino acid sequence selected from the group consisting of SEQ ID NO:2, SEQ ID NO:4 and SEQ ID NO:6.

One embodiment of the present invention relates to a substantially purified coccidian CKI protein comprising or consisting of the amino acid sequence disclosed in Figure 4 as SEQ ID NO:2. The amino acid sequence set forth in SEQ ID NO:2 represents the alpha ('α') isoform of a novel coccidian CKI identified from *Eimeria tenella*, designated EtCKIα. Another embodiment of the present invention relates to a substantially purified coccidian CKI protein

comprising or consisting of the amino acid sequence disclosed in Figure 4 as SEQ ID NO:4 or SEQ ID NO:6. The amino acid sequences set forth in SEQ ID NO:4 and SEQ ID NO:6 represent the alpha (' α ') and beta (' β ') isoforms of novel coccidian CKI identified from *Toxoplasma gondii*, designated TgCKI α and TgCKI β , respectively.

5 A preferred aspect of the present invention is disclosed in Figure 4 as SEQ ID NO:2, a coccidian amino acid sequence representing the *Eimeria tenella* casein kinase I alpha isoform, EtCKI α , and disclosed as follows:

1 MDVRVGKYYR LGRKIGSGSF GDIYLGNTIS TGDEVAIKLE SVRSRHPQLI YESKLYKILT
61 GGIGIPTLYW YGIEGDYNVM IIELLGPSLE DLFSICNRKL SLKTVLMLAD QMLNRIEFVH
10 121 SRHFHHRDIK PDNFLIGRGK KMSIVFAIDF GLAKKYRDPR TQSHIPYREG KNLGTGTARYA
181 SVNTHLGIEQ SRRDDLEALG YVLMYFNRRS LPWQGLKATT KDKYDKIME KKMSTPIEVL
241 CKQFPFEFIT YLNYCRSLRF EDRPDYSYLR RLFKDLFFRE GYQYDFIFDW TFLHAERERE
301 RQRRSMVNQG AESGNQWRRD ASGRDPLGRL PQLEP (SEQ ID NO:2).

15 Another preferred aspect of the present invention is disclosed in Figure 4 as SEQ ID NO:4, a coccidian amino acid sequence representing the *Toxoplasma gondii* casein kinase I alpha isoform, TgCKI α , and disclosed as follows:

1 MEVRVGKYYR LGRKIGSGSF GDIYIGANIL TGDEVAIKLE SIKSKHPQLL YESKLYKLLA
61 GGIGIPMVHW YGIEGDYNVM VIDLLGPSLE DLFSICNRKL SLKTVLMLAD QMLNRIEFVH
121 SKNFHHRDIK PDNFLIGRGK KMSVVYIIDF GLAKKYRDPK TQQHIPPYREG KNLGTGTARYA
20 181 SINTHLGIEQ SRRDDLEALG YVLMYFNRRS LPWQGLKATT KDKYDKIME KKMSTPIEIL
241 CKHFPFEFIT YLNYCRSLRF EDRPDYAYLR RLFKDLFFRE GYQYDFIFDW TFINTKEDRA
301 SRRSQVYVE DNRQVEENQN ELPM (SEQ ID NO:4).

25 Another preferred aspect of the present invention is disclosed in Figure 4 as SEQ ID NO:6, a coccidian amino acid sequence representing the *Toxoplasma gondii* casein kinase I beta isoform, TgCKI β , and disclosed as follows:

1 MAHQDTRNH TGVGPSSSIP LKDLKIAGVW KIGRKIGSGS FGDIYKGLNS QTGQEVALKV
61 ESTKAKHPQL LYEYKLLKHL QGGTGIAQVF CCETAGDHNI MAMELLGPSL EDVFNLCNRT
121 FSLKTILLLA DQFLQRVEYI HSKNFHHRDI KPDNLLGGA GNQNTIYVID FGLAKKFRDP
181 KTHQHIPPYRE NKNLTGTARY ASISAHLGSE QSRDDLEAV GYVLMYFCRG GTLPWQGIKA
30 241 NTKQEKYHKI MEKKMSTPVE VLCKGYPSEF ATYLHYCRSL RFEDRPDYAY LKRLFRDLYI
301 KEGYDDSDRE FDWTVKLSSR SLGPPSSRAQ HVLLSQDTRT RGKRETDPRV AARSGDRERG
361 IHFSNGNVGN PSMATNPGGL SVMVHERTSL VDQGDGRSRE TSTRKEDAKD GRWPGGRFSC
421 LPLLRRSPT KA (SEQ ID NO:6).

35 The present invention further relates to a substantially purified coccidian CKI protein, including but not limited to a CKI protein of the *Eimeria* or *Toxoplasma* genera, said

protein comprising at least about 80% amino acid sequence identity with an amino acid sequence selected from the group consisting of SEQ ID NO:2, SEQ ID NO:4 and SEQ ID NO:6.

Another embodiment of the present invention relates to a substantially purified coccidian CKI protein, including but not limited to a CKI protein of the *Eimeria* or *Toxoplasma* genera, said protein comprising at least about 80% amino acid sequence identity with an amino acid sequence selected from the group consisting of SEQ ID NO:2 and SEQ ID NO:4.

The present invention also relates to a substantially purified coccidian CKI protein, including but not limited to a CKI protein of the *Eimeria* or *Toxoplasma* genera, said protein comprising at least about 80% amino acid sequence identity with an amino acid sequence selected from the group consisting of SEQ ID NO:4 and SEQ ID NO:6.

The present invention further relates to a substantially purified coccidian CKI protein, including but not limited to a CKI protein of the *Eimeria* or *Toxoplasma* genera, said protein comprising at least about 80% amino acid sequence identity with an amino acid sequence as set forth in SEQ ID NO:2.

The present invention still further relates to a substantially purified coccidian CKI protein, including but not limited to a CKI protein of the *Eimeria* or *Toxoplasma* genera, said protein comprising at least about 80% amino acid sequence identity with an amino acid sequence as set forth in SEQ ID NO:4.

The present invention also relates to a substantially purified coccidian CKI protein, including but not limited to a CKI protein of the *Eimeria* or *Toxoplasma* genera, said protein comprising at least about 80% amino acid sequence identity with an amino acid sequence as set forth in SEQ ID NO:6.

The present invention also relates to biologically active fragments and/or mutants of the coccidian CKI proteins as initially set forth as SEQ ID NO:2, SEQ ID NO:4 and SEQ ID NO:6, including but not necessarily limited to amino acid substitutions, deletions, additions, amino terminal truncations and carboxy-terminal truncations such that these mutations provide for proteins or protein fragments of diagnostic, therapeutic or prophylactic use and would be useful for screening for agonists and/or antagonists for CKI function.

A preferred aspect of the present invention relates to a substantially purified, fully processed (including any proteolytic processing, glycosylation and/or phosphorylation) and mature coccidian CKI protein obtained from a recombinant host cell (both prokaryotic and eukaryotic, as well as both stably and transiently transformed/transfected cells) containing a DNA expression vector comprising a nucleotide sequence as set forth in SEQ ID NOs:1, 3 and/or 5, and expresses the *Eimeria* or *Toxoplasma* CKI precursor or mature form of the respective

protein. It is especially preferred that the recombinant host cells be a eukaryotic host cell, including but not limited to a mammalian or insect cell line.

The present invention also relates to subcellular membrane fractions and cell lysates of the recombinant host cells (both prokaryotic and eukaryotic, as well as both stably and transiently transformed/transfected cells) comprising the nucleic acid molecules of the present invention, including but not limited to the *Eimeria* and *Toxoplasma* CKI cDNA molecules disclosed herein. For a coccidian CKI protein that localizes to a membraneous region of the recombinant cells in which it is expressed, the subcellular membrane fractions can comprise wild-type or mutant forms of said coccidian CKI protein, often at levels substantially above wild-type levels and hence will be useful in various assays described throughout this specification. For a coccidian CKI protein that localizes to the cytosol of the recombinant cells in which it is expressed, a cell lysate fraction of said cells can comprise wild-type or mutant forms of said coccidian CKI protein. Again, the expression levels of said CKI protein can be substantially above wild-type levels and thus can be useful in various assays described herein.

Therefore, the present invention relates to methods of expressing the coccidian CKI genes and biological equivalents disclosed herein, assays employing these gene products, cells expressing these gene products, and agonistic and/or antagonistic compounds identified through the use of these coccidian CKI genes and expressed coccidian CKI protein, including, but not limited to one or more modulators of *Eimeria tenella* CKI α , *Toxoplasma gondii* CKI α and *Toxoplasma gondii* CKI β . These modulators may act through many different mechanisms, including but not limited to direct contact with the kinase domain of the coccidian CKI proteins or by preventing the binding of the target protein substrate to the coccidian CKI. Said modulators may either prevent or promote receptor activity.

As used herein, a "biologically active equivalent" or "functional derivative" of a wild-type coccidian CKI, including but not limited to a CKI protein of the *Eimeria* or *Toxoplasma* genera, possesses a biological activity that is substantially similar to the biological activity of the wild-type coccidian CKI. The term "functional derivative" is intended to include the "fragments," "mutants," "variants," "degenerate variants," "analogs," "homologues" or "chemical derivatives" of the wild-type coccidian CKI protein. The term "fragment" is meant to refer to any polypeptide subset of wild-type coccidian CKI. The term "mutant" is meant to refer to a molecule that may be substantially similar to the wild-type form but possesses distinguishing biological characteristics. Such altered characteristics include but are in no way limited to altered substrate binding, altered substrate affinity and altered sensitivity to chemical compounds affecting biological activity of the coccidian CKI or coccidian CKI functional derivative. The term "variant" is meant to refer to a molecule substantially similar in structure and function to

either the entire wild-type protein or to a fragment thereof. A molecule is "substantially similar" to a wild-type coccidian CKI-like protein if both molecules have substantially similar structures or if both molecules possess similar biological activity. Therefore, if the two molecules possess substantially similar activity, they are considered to be variants even if the structure of one of the molecules is not found in the other or even if the two amino acid sequences are not identical. The term "analog" refers to a molecule substantially similar in function to either the full-length coccidian CKI protein or to a biologically active fragment thereof.

Any of a variety of procedures may be used to clone a coccidian CKI of the present invention. These methods include, but are not limited to, (1) a RACE PCR cloning technique (Frohman, et al., 1988, *Proc. Natl. Acad. Sci. USA* 85: 8998-9002). 5' and/or 3' RACE may be performed to generate a full-length cDNA sequence. This strategy involves using gene-specific oligonucleotide primers for PCR amplification of coccidian CKI cDNA. These gene-specific primers are designed through identification of an expressed sequence tag ("EST") nucleotide sequence which has been identified by searching any number of publicly available nucleic acid and protein databases; (2) direct functional expression of the coccidian CKI cDNA following the construction of a coccidian CKI-containing cDNA library in an appropriate expression vector system; (3) screening a coccidian CKI-containing cDNA library constructed in a bacteriophage or plasmid shuttle vector with a labeled degenerate oligonucleotide probe designed from the amino acid sequence of the coccidian CKI protein; (4) screening a coccidian CKI-containing cDNA library constructed in a bacteriophage or plasmid shuttle vector with a partial cDNA encoding the coccidian CKI protein. This partial cDNA is obtained by the specific PCR amplification of coccidian CKI DNA fragments through the design of degenerate oligonucleotide primers from the amino acid sequence known for other kinases which are related to the coccidian CKI protein; (5) screening a coccidian CKI-containing cDNA library constructed in a bacteriophage or plasmid shuttle vector with a partial cDNA encoding, for example, an *Eimeria* or *Toxoplasma* CKI protein. This strategy may also involve using gene-specific oligonucleotide primers for PCR amplification of coccidian CKI cDNA identified as an EST as described above; or (6) designing 5' and 3' gene specific oligonucleotides using SEQ ID NOs:1, 3 or 5 as a template so that either the full-length cDNAs may be generated by known RACE techniques, or a portion of the coding regions may be generated by these same known RACE techniques to generate and isolate a portion of the coding regions to use as probes to screen one of numerous types of cDNA and/or genomic libraries in order to isolate a full-length version of the nucleotide sequence encoding coccidian CKI.

It is readily apparent to those skilled in the art that other types of libraries, as well as libraries constructed from other cell types or species types, may be useful for isolating a

coccidian CKI-encoding DNA, including but not limited to a CKI-encoding DNA from the *Eimeria* or *Toxoplasma* genus, or a coccidian CKI homologue. Other types of libraries include, but are not limited to, cDNA libraries derived from cells or cell lines other than coccidian cells or tissue such as a vertebrate host which may contain coccidian CKI-encoding DNA. Additionally, a coccidian CKI genes and homologues may be isolated by oligonucleotide- or polynucleotide-based hybridization screening of a coccidian genomic library. It is readily apparent to those skilled in the art that suitable cDNA libraries may be prepared from cells or cell lines which have CKI activity. Additionally, since the coccidian protozoa have multiple infectious stages (*e.g.*, sporozoites, merozoites, schizonts, oocysts for *Eimeria sp.*; sporozoites, tachyzoites, bradyzoites and oocysts for *Toxoplasma*), a cDNA library from which to clone the coccidian CKI genes of the present invention can be generated from any of these stages of the protozoan life-cycle. The selection of cells or cell lines for use in preparing a cDNA library to isolate a cDNA encoding a coccidian CKI may be done by first measuring cell-associated CKI activity using any known assay available for such a purpose.

Preparation of cDNA libraries can be performed by standard techniques well known in the art. Well known cDNA library construction techniques can be found for example, in Sambrook et al., 1989, *supra*. Complementary DNA libraries may also be obtained from numerous commercial sources, including but not limited to Clontech Laboratories, Inc. and Stratagene.

It is also readily apparent to those skilled in the art that DNA encoding coccidian CKI may also be isolated from a suitable genomic DNA library. Construction of genomic DNA libraries can be performed by standard techniques well known in the art. Well known genomic DNA library construction techniques can be found in Sambrook, et al., *supra*.

In order to clone a coccidian CKI genes of the present invention by one of the preferred methods, the amino acid or DNA sequences of a coccidian CKI, including but not limited to an *Eimeria* or *Toxoplasma* CKI protein, or a homologous protein, may be necessary. To accomplish this, the CKI protein or a protein homologue may be purified and partial amino acid sequence determined by automated sequenators. It is not necessary to determine the entire amino acid sequence, but the linear sequence of two regions of six (6) to eight (8) amino acids can be determined for PCR amplification of a partial coccidian CKI DNA fragment. Once suitable amino acid sequences have been identified, the DNA sequences capable of encoding them are synthesized. Because the genetic code is degenerate, more than one codon may be used to encode a particular amino acid, and therefore, the amino acid sequence can be encoded by any of a set of similar DNA oligonucleotides. Only one member of the set will be identical to the *T. gondii* and *E. tenella* CKI sequences disclosed herein, but others in the set will be capable of

hybridizing to these coccidian CKI DNAs even in the presence of DNA oligonucleotides with mismatches. The mismatched DNA oligonucleotides may still sufficiently hybridize to the specific coccidian CKI DNA to permit identification and isolation of coccidian CKI encoding DNA. Alternatively, the nucleotide sequence of a region of an expressed sequence may be identified by searching one or more available genomic databases. Gene-specific primers may be used to perform PCR amplification of a cDNA of interest from either a cDNA library or a population of cDNAs. As noted above, the appropriate nucleotide sequence for use in a PCR-based method may be obtained from SEQ ID NOs:1, 3 or 5, either for the purpose of isolating overlapping 5' and 3' RACE products for generation of a full-length sequence coding for coccidian CKI, or to isolate a portion of the nucleotide sequence coding for coccidian CKI for use as a probe to screen one or more cDNA- or genomic-based libraries to isolate a full-length sequence encoding coccidian CKI or coccidian CKI-like proteins.

It is also readily apparent to those skilled in the art that DNA encoding coccidian CKI, including but not limited to CKI from the *Eimeria* or *Toxoplasma* genera, may be synthetically generated. Many different methods are used for assembling and generating synthetic genes are known in the art. For example in one such method, a series of sequentially overlapping oligonucleotides are synthesized. The oligonucleotides anneal to form a double stranded DNA fragment containing nicks on both strands. DNA ligase, an enzyme that catalyses the formation of phosphodiester bonds between the 5'-phosphate of one double-strand oligonucleotide fragment and the 3'-hydroxyl terminus on another adjacent double-strand oligonucleotide, is used to seal the nicks. Synthetic genes can also be made using the template-directed and primer-dependent 5'- to 3'-synthesis capabilities of the large subunit of the enzyme DNA-Polymerase I (Klenow fragment). The polymerase uses deoxynucleoside-triphosphates to fill in gaps once end annealing of the long oligonucleotides occurs. Any nick in the resulting double-stranded DNA is sealed by DNA ligase. Finally, very long oligonucleotide chains can be synthesized so that their 3'-ends overlap upon annealing. A subsequent filling-in reaction using DNA polymerase completes the full-length, double-stranded DNA. A number of companies specialize in generating synthetic genes with a high degree of sequence accuracy including Entelechon GmbH (Regensburg, Germany) and MCLAB (South San Francisco, CA).

In an exemplified method, the *Toxoplasma* CKI enzymes of the present invention were identified based on EST sequence information mined from the *Toxoplasma* EST database (<http://www.toxodb.org/ToxoDB.shtml>; Li, A.L. et al., 2003, *Genome Res.* 13:443-454). Two distinct classes of RT-PCR product were amplified from tachyzoite RNA using PCR primers designed from a segment of overlapping EST clones that showed homology to CKI enzymes of other organisms (GenBank Accession Nos. BM175598 and T62400). The cloned RT-PCR

products were used to screen a tachyzoite cDNA library, and cDNAs corresponding to two different CKI genes were identified. Putative open reading frames were assigned based on sequence alignments of previously characterized CKI enzymes (Klimczak, L.J. et al., 1995, *Plant Physiol.* 109:687-696; Barik, S. et al., 1997, *supra*; Gross, S.D. and Anderson, R.A., 1998, *supra*; Moreno-Bueno, G. et al., 2000, *Biochem. J.* 349:527-537) and optimal translational start sites were identified. The two cDNAs identified from *T. gondii* have predicted amino acid products of 38 kDa and 49 kDa. Due to its higher degree of homology with *Plasmodium falciparum* CKI α (PfCKI α), the 38 kDa product was designated isoform " α " (TgCKI α), and the larger form isoform " β " (TgCKI β). TgCKI α shares 68% identity with PfCKI α at the amino acid level; while TgCKI β shares a 45% identity with the *Plasmodium* protein. The two CKI isoforms from *T. gondii* share 48% identity. An internal amino acid sequence of TgCKI α matches the TgCKI peptide identified in Knockaert et al., 2000, *supra*. The *Eimeria tenella* homologue of the *T. gondii* CKI α enzyme was cloned by screening a sporozoite cDNA library with a TgCKI α probe under low stringency conditions. Full-length clones encoding a predicted open-reading frame of 39 kDa were identified. Since the CKI isoform identified from *E. tenella* shares 81% identity at the amino acid level with TgCKI α , it was designated isoform " α " (EtCKI α).

Coccidian CKI cDNA, including but not limited to the *E. tenella* and *T. gondii* CKI genes disclosed herein, that are obtained through the methods described above may be recombinantly expressed by molecular cloning into an expression vector (such as pcDNA3.neo, pcDNA3.1, pCR2.1, pBlueBacHis2, pLITMUS28 or pETblue1 vector) containing a suitable promoter and other appropriate transcription regulatory elements, and transferred into prokaryotic or eukaryotic host cells to produce recombinant coccidian CKI. Expression vectors are defined herein as DNA sequences that are required for the transcription of cloned DNA and the translation of their mRNAs in an appropriate host. Such vectors can be used to express eukaryotic DNA in a variety of recombinant host cells such as bacteria, blue green algae, plant cells, protozoan cells, insect cells and mammalian cells. An appropriately constructed expression vector should contain the following: an origin of replication for autonomous replication in host cells, selectable markers, a limited number of useful restriction enzyme sites, a potential for high copy number, and active promoters. A promoter is defined as a DNA sequence that directs RNA polymerase to bind to DNA and initiate RNA synthesis. A strong promoter is one which causes mRNAs to be initiated at high frequency. Methods to determine the coccidian CKI cDNA sequence(s) that yields optimal levels of expressed coccidian CKI protein are well known in the art. Following determination of the coccidian CKI cDNA cassette yielding optimal expression, this coccidian CKI cDNA construct is transferred to a variety of expression vectors (including recombinant viruses), including but not limited to those for mammalian cells, plant cells, insect

cells, protozoan cells, oocytes, bacteria and yeast cells. Techniques for such manipulations can be found in Sambrook, et al., *supra*, and are well known and available to artisans of ordinary skill in the art. Therefore, another aspect of the present invention includes host cells that have been engineered to contain and/or express DNA sequences encoding coccidian CKI. An expression vector containing DNA encoding coccidian CKI protein may be used for expression of coccidian CKI in a recombinant host cell. Such recombinant host cells can be cultured under suitable conditions to produce coccidian CKI or a biologically equivalent form. Expression vectors may include, but are not limited to, cloning vectors, modified cloning vectors, specifically designed plasmids or viruses. Also, a variety of commercially available mammalian, bacterial, fungal cell, and insect cell expression vectors may be used to express recombinant coccidian CKI in the respective cell types, in addition to those expression vectors described in Donald, R.G.K. and Liberator, P.A., 2002 (*Mol. Biochem. Parasitol.* 120:165-175) and Donald, R.G.K. et al., 2002 (*Eukaryotic Cell.* 1:317-328).

Recombinant host cells may be prokaryotic or eukaryotic, including but not limited to, bacteria such as *E. coli*, fungal cells such as yeast, mammalian cells including, but not limited to, cell lines of bovine, porcine, monkey, and rodent origin; and insect cells. Protozoan host cells may also be used for transgenic expression of the CKI proteins disclosed herein, including but not limited to *Toxoplasma* and *Leishmanial* expression systems. Additionally, it may be beneficial that an inducible expression system (*e.g.*, tetracycline on/off system) be implemented for expression of said proteins.

The expression vector may be introduced into host cells via any one of a number of techniques including but not limited to transformation, transfection, protoplast fusion, and electroporation. The expression vector-containing cells are individually analyzed to determine whether they produce coccidian CKI protein. Identification of coccidian CKI expressing cells may be done by several means, including but not limited to immunological reactivity with anti-coccidian CKI antibodies, labeled ligand binding and the presence of host cell-associated coccidian CKI activity.

Expression of coccidian CKI DNA may also be performed using *in vitro* produced synthetic mRNA. Synthetic mRNA can be efficiently translated in various cell-free systems, including but not limited to wheat germ extracts and reticulocyte extracts, as well as efficiently translated in cell based systems, including but not limited to microinjection into frog oocytes, with microinjection into frog oocytes being preferred.

Levels of coccidian CKI in host cells is quantified by a variety of techniques, including but not limited to immunoaffinity and/or ligand affinity techniques. CKI-specific affinity beads or CKI-specific antibodies are used to isolate ³⁵S-methionine labeled or unlabelled

CKI. Labeled CKI protein is analyzed by SDS-PAGE. Unlabelled CKI protein is detected by Western blotting, ELISA or RIA assays employing either CKI protein specific antibodies and/or antiphosphothreonine/antiphosphoserine antibodies.

Following expression of CKI in a host cell, CKI protein may be recovered to provide CKI protein in active form. Several CKI protein purification procedures are available and suitable for use. Recombinant CKI protein may be purified from cell lysates and extracts, or from conditioned culture medium, by various combinations of, or individual application of salt fractionation, ion exchange chromatography, size exclusion chromatography, hydroxylapatite adsorption chromatography and hydrophobic interaction chromatography.

In addition, recombinant CKI protein can be separated from other cellular proteins by use of an immunoaffinity column made with monoclonal or polyclonal antibodies specific for full-length CKI protein or polypeptide fragments of CKI protein. Additionally, polyclonal or monoclonal antibodies may be raised against a synthetic peptide (usually from about 9 to about 25 amino acids in length) from a portion of the proteins as disclosed in SEQ ID NOs:2, 4 or 6. Monospecific antibodies to coccidian CKI are purified from mammalian antisera containing antibodies reactive against coccidian CKI or are prepared as monoclonal antibodies reactive with coccidian CKI using the technique of Kohler and Milstein (1975, *Nature* 256: 495-497).

Monospecific antibody as used herein is defined as a single antibody species or multiple antibody species with homogenous binding characteristics for coccidian CKI.

Homogenous binding, as used herein, refers to the ability of the antibody species to bind to a specific antigen or epitope, such as those associated with coccidian CKI, as described above. Coccidian CKI-specific antibodies are raised by immunizing animals such as mice, guinea pigs, rabbits, goats, horses and the like, with an appropriate concentration of coccidian CKI protein or a synthetic peptide generated from a portion of a coccidian CKI with or without an immune adjuvant. Preimmune serum is collected prior to the first immunization. Each animal receives between about 0.1 mg and about 1000 mg of coccidian CKI protein associated with an acceptable immune adjuvant, including but not limited to Freund's complete, Freund's incomplete, alum-precipitate, water in oil emulsion containing *Corynebacterium parvum* and tRNA. The initial immunization consists of coccidian CKI protein or a peptide fragment thereof in, preferably, Freund's complete adjuvant at multiple sites either subcutaneously (SC), intraperitoneally (IP) or both. The animals may or may not receive booster injections following the initial immunization depending on determination of antibody titer. At about 7 days after each booster immunization, or about weekly after a single immunization, the animals are bled, serum collected, and aliquots stored at about -20°C.

Monoclonal antibodies (mAb) reactive with coccidian CKI protein are prepared by immunizing inbred mice, preferably Balb/c, with coccidian CKI protein. The mice are immunized by the IP or SC route with about 1 mg to about 100 mg, preferably about 10 mg, of coccidian CKI protein in about 0.5 ml buffer or saline incorporated in an equal volume of an acceptable adjuvant, as discussed above. Immunized mice are given one or more booster immunizations by the intravenous (IV) route. Lymphocytes, from antibody positive mice, preferably splenic lymphocytes, are obtained by removing spleens from immunized mice by standard procedures known in the art. Hybridoma cells are produced by mixing the splenic lymphocytes with an appropriate fusion partner, preferably myeloma cells, under conditions which will allow the formation of stable hybridomas. The antibody producing cells and myeloma cells are fused in polyethylene glycol. Fused hybridoma cells are selected by growth in hypoxanthine, thymidine and aminopterin supplemented Dulbecco's Modified Eagles Medium (DMEM) by procedures known in the art. Supernatant fluids are collected from growth positive wells and are screened for antibody production by an immunoassay such as solid phase immunoradioassay (SPIRA) using coccidian CKI as the antigen. The culture fluids are also tested in the Ouchterlony precipitation assay to determine the isotype of the mAb. Hybridoma cells from antibody positive wells are cloned by a technique such as the soft agar technique of MacPherson, 1973, Soft Agar Techniques, in *Tissue Culture Methods and Applications*, Kruse and Paterson, Eds., Academic Press.

Monoclonal antibodies are produced *in vivo* by injection of pristine primed Balb/c mice, approximately 0.5 ml per mouse, with about 2×10^6 to about 6×10^6 hybridoma cells about 4 days after priming. Ascites fluid is collected at approximately 8-12 days after cell transfer and the monoclonal antibodies are purified by techniques known in the art.

In vitro production of anti-coccidian CKI mAb is carried out by growing the hybridoma in DMEM containing about 2% fetal calf serum to obtain sufficient quantities of the specific mAb. The mAb are purified by techniques known in the art.

Antibody titers of ascites or hybridoma culture fluids are determined by various serological or immunological assays known in the art. Similar assays are used to detect the presence of coccidian CKI in fluids or tissue and cell extracts.

It is readily apparent to those skilled in the art that the above described methods for producing monospecific antibodies may be utilized to produce antibodies specific for a coccidian CKI peptide fragments, or a respective a full-length coccidian CKI.

The coccidian CKI proteins of the present invention are suitable for use in an assay procedure for the identification of compounds which modulate CKI activity. A CKI-containing fusion construct, such as a FLAG-epitope tagged CKI fusion protein as discussed

within this specification, is useful to measure CKI activity. Kinase activity can be measured, for example, using a modified version of the casein kinase I assay described by Donald et al., R.G.K. et al., 2002, *supra*, and as outlined in Example 2. Casein kinase activity can be assayed using 96-well phosphocellulose plates to capture ^{33}P -phosphorylated peptide or casein substrates. Kinase reactions can be carried out in 40 μl volumes containing 25 mM Hepes pH 7.4, 10 mM MgCl_2 , 20 mM β -glycerophosphate, 1 mM DTT, 250 μM substrate, 1 mg/ml BSA, 2 μM ATP and 0.01 μM [γ - ^{33}P]ATP. The reaction is initiated by addition of enzyme and then incubated at room temperature for 60 minutes. A variety of commercially available peptides substrates can be use, as described in Example 2.

The coccidian CKI proteins of the present invention, including but not limited to CKI proteins of the *Eimeria* and *Toxoplasma* genera, may be obtained from both native and recombinant sources (as a full-length protein, biologically active protein fragment, or fusion construction) for use in assay procedures to identify coccidian CKI modulators. Modulating CKI includes the inhibition or activation of the kinase, representing antagonists or agonists, respectively. In general, an assay procedure to identify coccidian CKI modulators will contain at least a functional domain of a coccidian CKI and a test compound or sample which contains a putative CKI kinase agonist or antagonist. The test compound or sample may be tested directly on, for example, purified CKI or an epitope tagged-CKI fusion, subcellular fractions of CKI-producing cells containing CKI (native or recombinant), whole cells expressing coccidian CKI (native or recombinant), and/or CKI protein fragments and respective deletion fragments. The test compound or sample may be added to CKI in the presence or absence of a known CKI substrate, including but not limited to casein. The modulating activity of the test compound or sample may be determined by, for example, analyzing the ability of the test compound or sample to bind to CKI, activate the protein, inhibit the protein, inhibit or enhance the binding of other compounds to coccidian CKI, or modifying kinase activity.

The present invention is also directed to methods for screening for compounds which modulate the expression of DNA or RNA encoding a coccidian CKI protein, including but not limited to a CKI protein of the *Eimeria* or *Toxoplasma* genera. Compounds which modulate these activities may be DNA, RNA, peptides, proteins, or non-proteinaceous organic molecules. Compounds may modulate CKI protein by increasing or attenuating the expression of DNA or RNA encoding the protein or the function of coccidian CKI. Compounds that modulate the expression of DNA or RNA encoding coccidian CKI or the biological function thereof may be detected by a variety of assays. The assay may be a simple "yes/no" assay to determine whether there is a change in expression or function. The assay may be made quantitative by comparing the expression or function of a test sample with the levels of expression or function in a standard

sample. Kits containing at CKI, antibodies to coccidian CKI, or modified coccidian CKI may be prepared by known methods for such uses.

The DNA molecules, RNA molecules, recombinant proteins and antibodies of the present invention may be used to screen and measure levels of coccidian CKI. The recombinant proteins, DNA molecules, RNA molecules and antibodies lend themselves to the formulation of kits suitable for the detection and typing of coccidian CKI. Such a kit would comprise a compartmentalized carrier suitable to hold in close confinement at least one container. The carrier would further comprise reagents such as recombinant CKI or anti-CKI antibodies suitable for detecting coccidian CKI. The carrier may also contain a means for detection such as labeled antigen or enzyme substrates or the like.

Pharmaceutically useful compositions comprising modulators of coccidian CKI may be formulated according to known methods such as by the admixture of a pharmaceutically acceptable carrier. Examples of such carriers and methods of formulation may be found in Remington's Pharmaceutical Sciences. To form a pharmaceutically acceptable composition suitable for effective administration, such compositions will contain an effective amount of the protein, DNA, RNA, modified coccidian CKI, or either CKI agonists or antagonists including serine or threonine kinase activators or inhibitors.

Therapeutic or diagnostic compositions of the invention are administered to an animal, including but not limited to humans and poultry, in amounts sufficient to treat or diagnose disorders. The effective amount may vary according to a variety of factors such as the animal's condition, weight, sex and age. Other factors include the mode of administration.

The pharmaceutical compositions may be provided to the individual by a variety of routes such as subcutaneous, topical, oral and intramuscular.

The term "chemical derivative" describes a molecule that contains additional chemical moieties which are not normally a part of the base molecule. Such moieties may improve the solubility, half-life, absorption, etc. of the base molecule. Alternatively the moieties may attenuate undesirable side effects of the base molecule or decrease the toxicity of the base molecule. Examples of such moieties are described in a variety of texts, such as Remington's Pharmaceutical Sciences.

All publications mentioned herein are incorporated by reference for the purpose of describing and disclosing methodologies and materials that might be used in connection with the present invention. Nothing herein is to be construed as an admission that the invention is not entitled to antedate such disclosure by virtue of prior invention.

Having described preferred embodiments of the invention with reference to the accompanying figures, it is to be understood that the invention is not limited to those precise

embodiments, and that various changes and modifications may be effected therein by one skilled in the art without departing from the scope or spirit of the invention as defined in the appended claims.

The following examples are provided to illustrate the present invention without, however, limiting the same hereto.

EXAMPLE 1

Identification of cDNAs Encoding Coccidian Parasite Casein Kinase I

Cloning of Toxoplasma gondii CKI α and CKI β - BLAST searching of the *Toxoplasma* EST database (<http://www.toxodb.org/ToxoDB.shtml>) identified a pair of overlapping ESTs (GenBank BM175598 and T62400) with homology to a CKI consensus sequence derived from a ClustalW alignment of CKI enzymes characterized from a variety of organisms (AlignX, Vector NTI suite). The *Toxoplasma* EST sequences identified are listed below:

BM175598

AGAATTTCAATTCACAGAGATATCAAACCAGATAACTTTCTTCTCGGCGGTGCCGGCAATCAAAACACGATCTACGTG
ATCGACTTCGGCCTGGCGAAGAAGTTTCGCGATCCGAAAACGCACCAACATATTCCGTACAGAGAAAACAAGAATCT
CACGGGAACGGCGCGCTACGCGTCCATCAGTGC GCATCTGTGTTCGAGCAGAGTCGCCGAGATGACCTCGAAGCAG
TCGGCTACGTTCTCATGTACTTCTGTTCGAGGAGGCACGCTGCCTTGGCAGGGCATCAAAGCGAATACCAAACAGGAG
AAGTACCACAAGATCATGGAGAAGAAGATGTTCGACGCCCCGTCGAGGTGCTATGCAAGGGATATCCAAGCGAATTTGC
CACATACTTGCACTACTGCCGCTCCTTGCGATTTCGAGGACCGACCGGACTACGCCTACCTCAAGCGACTCTTTTCGAG
ATCTCTACATCAAAGAGGGCTACGATGACAGTGACCGCGAATTCGACTGGACAGTGAACTTTCGTCGCGCAGTCTC
GGAC (SEQ ID NO:7).

T62400

GATATCCAAGCGAATTTGCCACATACTTGCACTACTGCCGCTCCTTGCGATTTCGAGGACCGACCGGACTACGCCTAC
CTCAAGCGACTCTTTTCGAGATCTCTACATCAAAGAGGGCTACGATGACAGTGACCGCGAATTCGACTGGACAGTGAA
ACTTTTCGTCGCGCAGTCTCGGACCGCCAAGCAGTCGAGCGCAACATGTTTACTGAGTCAAGACACCCGAACGCGAG
GGAAGCGGGAGACAGATCGACCTGTCTGCTGTGCGGAGTGGCGACCGCGAACGAGGAATCCATTTACGCAACGGGAAC
GTGGGCAATCCCTCCGATGGCAACGAACCCCCG (SEQ ID NO:8).

A series of nested PCR primers was designed and used to amplify cDNA fragments from tachyzoite RNA by RT-PCR. RNA was prepared with an Oligo (dT)₂₅

Dynabeads® kit (Dyna; Lake Success, NY) and a first-strand cDNA pool synthesized with superscript II (Invitrogen; Carlsbad, CA). PCR products were cloned with a TA cloning kit (Promega; Madison, WI) and sequenced. Most of the PCR products obtained corresponded to the original EST sequence. However, one of the three sets of PCR primer pairs used,

- 5 GATATCAAACCAGATAACTTTCTTCTCGGC (SEQ ID NO:9) and
 CAAGGAGCGGCAGTAGTGCAAGT (SEQ ID NO:10), also amplified a second class of
 cDNA product which showed greater homology to the CKI- α isoform of *Plasmodium falciparum*
 (Barik, S. et al., 1997, *supra*). The two distinct cDNA fragments were used separately to probe a
 tachyzoite cDNA library (#1896, NIH AIDS Research and Reference Reagent Program).
 10 Putative open reading frames were assigned based on sequence alignments of previously
 characterized CKI enzymes (Klimczak, L.J. et al., 1995, *supra*; Barik et al., 1997, *supra*; Gross
 and Anderson, 1998, *supra*; Moreno-Bueno, G. et al., 2000, *supra*), and the presence of an
 optimal nucleotide context surrounding potential translational start sites was identified (Seeber,
 F., 1997, *Parasitol. Res.* 83:309-311). Corresponding full-length clones were obtained and
 15 designated " α " or " β " based on their relative homology to malaria isoform PfCKI α . At least four
 different full-length clones with 5' untranslated nucleotide sequences extending beyond the
 predicted start site were obtained for each isoform. No cDNAs corresponding to potentially
 functional splice variants were detected. Since corresponding TgCKI " α " or " β " genomic contigs
 identified in the *Toxoplasma* genome database do not overlap
 20 (<http://www.toxodb.org/ToxoDB.shtml>), the genes appear to map to separate loci.

- Cloning of Eimeria tenella CKI α* - The *E. tenella* homologue of the *T. gondii* CKI
 alpha enzyme was cloned by screening a sporozoite cDNA library with a TgCKI α probe under
 low stringency conditions (plaque lifts hybridized in 30% formamide and washed with 5XSSPE,
 0.1% SDS at 42°C). The cDNA library was generated from polyA+ RNA purified from *Eimeria*
 25 sporozoites using a standard lambda ZAP cloning kit and packaging reagents (Stratagene; La
 Jolla, CA). The TgCKI α probe, corresponding to the TgCKI α open reading frame, was
 generated by PCR using the following primer set:
 ATGGACTACAAAGACGATGACGACAAGGAGGTCAGGGTCGGAGGCAAGTACCGAC
 (SEQ ID NO:24) and CGGTCTAGATCAGAGGGAGACGCGCGTCCTGACC (SEQ ID
 30 NO:25).

- Results* - Two *T. gondii* cDNAs encoding CKI isoforms have been identified,
 TgCKI α (Figure 2; SEQ ID NO:3) and TgCKI β (Figure 3; SEQ ID NO:5). These cDNAs
 encode protein products with predicted molecular weights of 38 kDa (Figure 4; SEQ ID NO:4)
 and 49 kDa (Figure 4, SEQ ID NO:6), respectively. Due to its higher degree of homology with
 35 the *Plasmodium falciparum* CKI α isoform (68% identify, Figure 5), the smaller protein was

designated isoform " α " (TgCKI α). An internal amino acid sequence of this isoform (shown in Figure 4) matches a TgCKI peptide that was identified by microsequencing of a 38 kDa protein with strong affinity for an immobilized CDK inhibitor (purvalanol B) matrix (Knockaert, M. et al., 2000, *Chem. Biol.* 7:411-422). CKI isoforms from *Plasmodium falciparum* (PfCKI α),
 5 *Leishmania mexicana* (LmCKI-2) and *Trypanosoma cruzi* (TcCKI-1) were also found tightly associated with this matrix (identified peptides also highlighted), and their sequences are included for comparison in Figure 4. Examination of the percentage identity matrix derived from pairwise sequence alignments (Figure 5) reveals that TgCKI α shares a greater degree of identity with these parasite CKI orthologues than does the larger *T. gondii* isoform. A cDNA encoding
 10 the homologue of TgCKI α has been identified from the coccidia *Eimeria tenella*, EtCKI α (Figure 1; SEQ ID NO:1), generating a protein product with a predicted molecular weight of 39 kDa (Figure 4; SEQ ID NO:2). The EtCKI α enzyme shares 81% identity with the TgCKI α homologue but only 48% identity with the TgCKI β isoform (Figure 5).

EXAMPLE 2

Transient Expression of Coccidian Casein Kinase I Isoforms in *T. gondii*

Epitope tagged constructs - N-terminal FLAG epitope tags were appended to TgCKI α and TgCKI β open reading frames by PCR amplification (KOD polymerase, Novagen; Madison, WI). The sense and antisense PCR primers used to amplify the open frames are as
 20 follows:
 ATGGACTACAAAGACGATGACGACAAGGAGGTCAGGGTCGGAGGCAAGTACCGAC (SEQ ID NO:24) and CGGTCTAGATCAGAGGGAGACGCGCGTCCTGACC (SEQ ID NO:25) for TgCKI α ; and
 25 ATGGACTACAAAGACGATGACGACAAGGCGCACCATCAAGACACCCGCAAC (SEQ ID NO:26) and CGGTCTAGATCAAAAAAAGAACTTACGCCCACGGCGT (SEQ ID NO:27) for TgCKI β . The resulting DNA fragments were subcloned into *E. coli* pETblue1 (Novagen). For subcloning into *T. gondii* tubulin-promoter expression vectors (Donald, R.G.K. and Liberator, P.A., 2002, *Mol. Biochem. Parasitol.* 120:165-175), sense primers bearing additional
 30 5' compatible restriction enzyme recognition sites (*Bam*HI and *Bcl*II) were used:
 GGCGGATCCGAAAATGGACTACAAAGACGATGACGACAAGGAGGTCAGGGTCGGA
 GGCAAGTACCGAC (SEQ ID NO:28) for TgCKI α ; and
 GGCGTGATCAAAAATGGACTACAAAGACGATGACGACAAGGCGCACCATCAAGAC
 ACCCGCAAC (SEQ ID NO:29) for TgCKI β .

Chimeric TgCKI β -CAT constructs used for the deletion mapping of the TgCKI β C-terminal domain were made by inserting PCR-amplified TgCKI β fragments into CAT fusion vector p_{ubXho}/CAT (Donald and Liberator, 2002, *supra*). PCR fragments bearing successive 3' end truncations were amplified from TgCKI β cDNA with KOD polymerase and featured flanking *Bcl*I and *Sall*/*Xho*I sites (5' and 3' ends, respectively) to facilitate directional sub-cloning. The 5' sense primer used to amplify said fragments is as follows:

GGCGTGATCAAAAATGGACTACAAAGACGATGACGACAAGGCGCACCATCAAGACACCCGCAAC (SEQ ID NO:29). The antisense 3' primers used to amplify said fragments are as follows: GGCCTCGAGGGCCTTCGTCGGAGAGCGCCGACATAACAGTG (SEQ ID NO:30) for the full-length; GGCGTCGACGATGTTATGGTCGCCCCGAGTCTCGCAACA (SEQ ID NO:31) for 3' Δ 332; GGCGTCGACGATCTTGTGGTACTTCTCCTGTTTGGTATTCGCTTTGATGC (SEQ ID NO:32) for 3' Δ 182; and GGCCTCGAGCACGTTCCCGTTGCTGAAATGGATTCTCCTCGTTC (SEQ ID NO:33) for 3' Δ 64 (see also Fig. 7).

The FLAG-epitope tagged EtCKI α open-reading frame was cloned into *E. coli* and *T. gondii* expression vectors using the same strategy as that outlined above for TgCKI enzymes. The following sense and antisense primers were used for PCR amplification and subsequent *E. coli* expression vector cloning:

ATGGACTACAAAGACGATGACGACAAGGACGTCCGTGTGGGGGGTAAGTATCGTTTG (SEQ ID NO:34) and CGGTCTAGATCACGGTTCTAACTGAGGCAACCGTCCAAGT (SEQ ID NO:35). To facilitate restriction enzyme mediated cloning into the Toxoplasma expression vector, the sense primer was modified to contain a 5' flanking *Bgl*II site: GGCAGATCTGAAAATGGACTACAAAGACGATGACGACAAGGACGTCCGTGTGGGGGGTAAGTATCGTTTG (SEQ ID NO:36).

Host cells and parasite cultures – RH strain or RH Δ HXGPRT strain tachyzoites of *T. gondii* were maintained by serial passage in confluent monolayers of human foreskin fibroblasts ("HFF") as described (Roos, D.S. et al., 1994, *Methods Cell Biol.* 45:27-63). A modified [³H]-uracil uptake assay (Pfefferkorn, E.R. and Pfefferkorn, L.C., 1977, *J. Protozool.* 24:449-453) adapted for use in 96-well scintillation plates (Cytostar-T, Amersham; Piscataway, NJ) was used to measure parasite growth inhibition. Each well, previously seeded with HFF cells, was inoculated with *T. gondii* tachyzoites (2×10^4) in 200 μ l of growth media containing 2 μ Ci [5,6]-³H-uracil (Perkin-Elmer NEN; Boston, MA) and serially diluted compound. Following incubation at 37°C for 36-48 hours, tritium-incorporation into host cell monolayers was directly counted in a microplate scintillation counter (MicroBeta, PerkinElmer-Wallac).

Transfection protocol – Tachyzoites were transfected with 100 μ g of expression

plasmid FLAG-TgCKI α , FLAG-TgCKI β , FLAG-EtCKI α or vector alone, in triplicate, and inoculated into cultures of HFF cell monolayers (slides or T-25 flasks).

Immunological reagents and techniques – *T. gondii* CKI epitopes used to raise antisera are shown in Figure 4. A C-terminal cysteine residue was added to each peptide (synthesized by SynPep; Dublin, CA) to facilitate coupling to keyhole-limpet-hemocyanin carrier protein (KLH) by maleimide chemistry (immunogen conjugation kit, Pierce; Rockford, IL). Antisera was generated against the coupled peptide-KLH immunogen (Covance; Denver, PA) and subsequently affinity-purified by peptide affinity chromatography (SulfoLink kit, Pierce). Antisera was similarly prepared against coccidian parasite calmodulin domain like protein kinase (CDPK1) (Kieschnick, H. et al., 2001, *J. Biol. Chem.* 276:12369-12377; Dunn, P.P. et al., 1996, *Parasitology* 113:439-448). Epitopes corresponding to the CDPK1 enzyme orthologue from *Eimeria tenella* were chosen for antisera production: AKDLIRKMLAYVPSMRISARD (SEQ ID NO:11) and AVKVISKRQVKQKTDKELLL (SEQ ID NO:12). The resulting affinity purified antisera recognize both the *Toxoplasma* CDPK1 and the *Eimeria* CDPK homologue (not shown).

At 24 hours post-infection, anti-FLAG indirect immunofluorescence analysis ("IFA") was performed. For IFA staining, HFF host cell cultures were grown on Falcon culture slides, infected with parasites and examined 24-36h later. Infected monolayers were fixed in 3% paraformaldehyde in PBS, permeabilized with 0.25% Triton-X100 (in PBS) and pre-treated with blocking buffer (1% BSA, 5% serum in PBS) before staining. Treated coverslips were examined with a Zeiss Axiovert 35 inverted microscope equipped with a 100W Hg-vapor lamp and FITC filter. Polyclonal antisera to the bacterial chloramphenicol acetyltransferase (CAT) reporter and FLAG epitope monoclonal antisera were obtained from Sigma (St. Louis, MO).

Kinases transiently expressed in *T. gondii* were immunoprecipitated with agarose beads conjugated with monoclonal FLAG antisera ("FLAG-beads") (TgCKI and EtCKI) or protein A-beads (TgCKI-CAT chimeras) and assayed as previously described (Donald and Liberator, 2002, *supra*).

Western blots were performed as per standard SDS-PAGE techniques, followed by electrophoretic transfer to nitrocellulose. The antiserum used to probe the nitrocellulose blot was the polyclonal antiserum specific for the CKI α -IT epitope which is well conserved among CKI enzymes (see Figure 4).

Protein kinase assays - Casein kinase activity was assayed essentially as described (Donald, R.G.K. et al., 2002, *supra*) using 96-well phosphocellulose plates (MAPH-NOB, Millipore; Billerica, MA) to capture ³³P-phosphorylated peptide or casein substrates. Kinase reactions were carried out in 40 μ l volumes containing 25 mM Hepes, pH 7.4, 10 mM MgCl₂, 20 mM β -glycerophosphate, 1 mM DTT, 250 μ M substrate, 1 mg/mL BSA, 2 μ M ATP, and

0.01 μM [γ - ^{33}P]ATP (Perkin-Elmer, 3000 Ci/mmol). The reactions were initiated by the addition of enzyme and incubated at room temperature for 60 min. Casein (5% solution), partially dephosphorylated α -casein and β -casein were obtained from Sigma.

A variety of commercially available peptide substrates were evaluated. From Calbiochem (San Diego, CA): CKI substrate (RRKDLHDDEEDEAMSITA; SEQ ID NO:13); CKII substrate (RRADDSDDDDD; SEQ ID NO:14); Syntide-2 substrate (PLARTLSVAGLPQKK; SEQ ID NO:15); CaM kinase II substrate (281-291) (MHKNETVECLK; SEQ ID NO:16). From Promega: CKI substrate (DDDEESITRR; SEQ ID NO:17) and kemptide (LRRASLG; SEQ ID NO:18). From New England Biolabs (Beverly, MA): CKI phosphopeptide substrate (KRRRALS(p)VASLPGL; SEQ ID NO:19), and CKII substrate (RRREEETEEE; SEQ ID NO:20). PKG assays were performed with kemptide as substrate and 10 μM cGMP. CDPK was assayed with Syntide-2 substrate and in the presence of 100 μM CaCl_2 .

Results – Plasmid constructs encoding N-terminal FLAG epitope tagged TgCKI α and TgCKI β open reading frames were transfected into tachyzoites and expression was monitored by indirect immunofluorescence analysis (IFA) following inoculation of parasites onto HFF cell monolayers (Figure 6A). Uniform staining consistent with cytosolic expression was found in intracellular tachyzoites transfected with the FLAGTgCKI α construct. In contrast, the FLAGTgCKI β plasmid yielded a surface membrane-associated staining pattern similar to that observed with plasmids expressing dually acylated forms of PKG (Donald and Liberator, 2002, *supra*). Flag epitope-tagged EtCKI α constructs were also transiently expressed in *T. gondii* tachyzoites (Figure 6A). Recombinant FLAGEtCKI α displayed a cytosolic expression pattern, similar to recombinant FLAGTgCKI α . As expected, Western blotting of immunoprecipitated material with antisera to a conserved internal CKI epitope detected the presence of a 38 kDa protein in transfections with the TgCKI α or the EtCKI α plasmids (Figure 6C). In the case of the TgCKI β construct, a 55 kDa band was observed, a mobility somewhat slower than predicted from its 49 kDa mass.

Significant levels of immunoprecipitated CKI activity were detected in transfections with TgCKI α and TgCKI β plasmids using milk casein as substrate (Figure 6B). Approximately two-fold higher levels of CKI activity were detected with the TgCKI α plasmid than with the TgCKI β construct, which in turn was eight-fold above background levels observed in mock transfections (Figure 6B). Immunoprecipitated casein kinase I activity was also detected in lysates of parasites transfected with the EtCKI α plasmid (Figure 6B), while no kinase activity was detected in mock-transfected parasites. As observed with TgCKI expression constructs, ectopic expression of EtCKI α was toxic to *T. gondii* tachyzoites. Parasites expressing high

levels of recombinant *T. gondii* or *E. tenella* CKI enzyme appeared sick (Figure 6A).

Due to the lack of obvious N-terminally located acylation signals in TgCKI β , the location of the determinant responsible for its membrane association was mapped. A TgCKI β - chloramphenicol acetyl transferase (TgCKI β -CAT) chimera was constructed that yielded qualitatively similar levels of membrane-associated CKI activity in transiently transfected parasites as the parental FLAGTgCKI β construct (Figure 7). In this case, kinase activity was immunoprecipitated, and subcellular location determined by IFA with antisera to the CAT reporter. Successive 3' TgCKI β deletions were made in the chimeric protein and the constructs similarly assessed in transient assays. Deletion of the C-terminal 64 amino acids of TgCKI β had no effect on CKI activity but abolished its ability to associate with the plasma membrane. This deletion (3' Δ 64-CAT) generated the same cytosolic staining observed with the FLAGTgCKI α plasmid. Tachyzoites transfected with the chimeric constructs were also subjected to selection in serial passages of 20 μ M chloramphenicol. Stable transgenic lines expressing protein chimeras that lack CKI activity were readily obtained, such as those missing the C-terminal 182 or 332 amino acids (3' Δ 332-CAT and 3' Δ 182-CAT, Figure 7). In contrast, no stable lines were obtained from transfections with chimeras exhibiting CKI-activity (TgCKI β -CAT and 3' Δ 64-CAT, Figure 7).

EXAMPLE 3

Purification and Characterization of the Coccidian CKI Isoforms

Purification of recombinant CKI enzymes – *E. coli* strain BL21 OrigamiTM (Novagen) was transformed with FLAGTgCKI α and FLAGTgCKI β expression vectors, and colonies were expanded into 250 mL LB cultures grown at 37°C. Expression was induced with 1 mM IPTG once cultures had reached an OD_{600nm} of 0.3-0.5, and the transformed bacteria were harvested 4-6 hours later. Cell pellets were resuspended in 15 mL of lysis buffer containing protease inhibitors and lysates prepared as described by the supplier (BugbusterTM reagents, Novagen). The concentration of NaCl in the lysate was adjusted to 0.5 M, and 0.5 mL of FLAG-agarose beads were added (Sigma). The mixture was allowed to rock in a Nutator mixer (Becton-Dickinson; San Jose, CA) for 2 hours at 4°C. The slurry was then poured into an empty column and the matrix washed with 50 mL of wash buffer (50mM Tris-HCl, pH 7.4, 0.5M NaCl, 10% glycerol, 1 mM DTT, 1 mM EDTA). Enzyme was eluted with 5 mL of 0.4 mg/mL FLAG peptide (Sigma) in NaCl-free wash buffer and concentrated with a centrifuge filtration unit (Amicon Ultra®, 10 kDa cut-off, Millipore). Protein was measured with a micro-BCA reagent (Pierce) following a TCA-sodium deoxycholate step to remove residual FLAG peptide (Brown et

al., 1989, *Anal. Biochem.* 180:136-139). Using this procedure, and SDS-PAGE analysis, the yield of FLAG-TgCKI β was estimated as ~200 μ g of enzyme with >90% purity. Purification of FLAG-TgCKI α was much less efficient with only about 1 μ g of ~10% pure enzyme obtained per 250 mL bacterial culture.

Preparations of FLAG-TgCKI α and FLAG-TgCKI β were purified further on a SMART system FPLC 0.1 mL MonoQ anion exchange column (Pharmacia; New York, NY) equilibrated in buffer A (50mM Hepes, pH 7.4, 10% glycerol, 1 mM DTT) at room temperature. Samples of FLAG-affinity purified enzyme (10-100 μ g) were syringe-passed through a 0.22 μ m (pore-size) filter before loading on the column. A linear salt gradient was applied (0-500 mM in 3 mL at 100 μ l/min) and 100 μ l fractions were collected and analyzed by SDS-PAGE and for CKI activity. Fractions with peak activity were retained for further analysis. The degree of purity of the partially purified FLAG-TgCKI α was estimated by densitometry of silver stained SDS-PAGE gels. A recombinant rat CKI δ enzyme used for control purposes was obtained from New England Biolabs.

Partial purification of native TgCKI α - Parasites from a total of 75X T175 flasks were filtered through 3 μ m (pore-size) polycarbonate membranes (Millipore) to remove host cell debris. Following centrifugation in 1 L bottles at 3,000xg for 15 minutes, cell pellets were washed in PBS, the volumes combined and re-centrifuged. The final cell pellet was gently resuspended in 10 mL of lysis buffer which consisted of buffer A supplemented with 1% NP40 and a protease inhibitor cocktail (CompleteTM; Boehringer, Ingelheim, Germany). The suspension was sonicated with a Branson Sonifier microtip, centrifuged at 100,000xg (30 min at 4°C), and the supernatant passed through a 0.22 μ m filter (Millipore). Approximately 9 mL was loaded onto a LKB FPLC System 5-mL HiTrapQ anion-exchange column (Pharmacia) and run at 4°C. A segmented NaCl gradient was applied (0-0.2 M in 60 mL, 0.2-0.5 M in 45 mL and 0.5-1.0 M in 0.1 mL at 5 mL/min flow rate). Fractions of 5 mL were collected and assayed for CKI activity.

Fraction 7 from the HiTrapQ run, which contained peak TgCKI α activity, was concentrated (Amicon Ultra® unit), filtered (0.22 μ m pore size), and approximately 0.5 mL was loaded onto a SMART system FPLC System 0.1-mL phenyl superose PC 1.6/5 hydrophobic interaction chromatography column (Pharmacia). A linear (NH₄)₂SO₄ gradient was applied (1.0 M-0 in 1.5 mL at 0.05 mL/min flow rate). Fractions of 0.1 mL were collected and assayed for CKI activity, and the active fractions corresponding to the single peak were pooled for serological and enzymatic analysis.

Partial purification of native EtCKI α - The inability to stably express recombinant EtCKI α in *T. gondii* provided impetus to purify native enzyme from *E. tenella*

parasites (slide 5). A lysate was prepared from 2×10^{10} unsporulated oocysts using previously published procedures (Gurnett, A.M. et al., 2002, *supra*). The first two steps of the purification scheme used were essentially scaled-up from the method described for the purification of native TgCKI α . The lysate was subjected to preparative HiLoad Q anion-exchange chromatography and fractions eluted with a NaCl gradient (0-1 M NaCl). Fractions with peak kinase activity were pooled and subjected to hydrophobic interaction chromatography and fractions were eluted with a linear (NH₄)₂SO₄ gradient (1 M to 0). A third and final cation exchange chromatography purification step resulted in active fractions of sufficient purity.

Compounds - A collection of ATP-site competitive inhibitors of CDKs were used (Knockaert, M. et al., 2002, *Trends Pharmacol. Sci.* 23:417-425). All but aminopurvalanol are currently available commercially (e.g., Alexis Corporation, Tocris, Calbiochem, and A.G. Scientific). They were stored as 10 mM stock solutions in DMSO.

Results - In order to obtain sufficient quantity and purity of TgCKI enzyme for biochemical characterization, recombinant FLAGTgCKI α and FLAGTgCKI β proteins were expressed in *E. coli* and purified in a two step procedure. After an initial FLAG-affinity purification step, partially purified FLAGTgCKI α and FLAGTgCKI β enzymes were subjected to anion exchange FPLC chromatography using a linear salt gradient to elute CKI activity. Fractions with peak CKI activity were analyzed by SDS-PAGE and stained with silver (Figure 8A). The purity of recombinant TgCKI α and TgCKI β enzymes was estimated by densitometry as 30%, and greater than 95%, respectively. The lower purity of the FLAGTgCKI α enzyme is a reflection of the relatively poor yield of soluble enzyme recoverable from *E. coli* lysates compared with the TgCKI β isoform. The identity and integrity of the recombinant enzymes was confirmed in Western blots (Figure 8B) with antisera raised against conserved and isoform selective epitopes (see Figure 4). As with recombinant enzymes expressed transiently in *T. gondii*, TgCKI α was observed as a 38 kDa polypeptide, while TgCKI β exhibited the mobility of a 55 kDa protein rather than the 49 kDa predicted from conceptual translation.

To investigate the activity of the corresponding native TgCKI enzymes in tachyzoites, a detergent extract derived from 2×10^{10} parasites was fractionated by HiTrapQ anion exchange chromatography (Figure 9A&B). Fractions eluted with a salt gradient were assayed for casein kinase activity using a phosphorylated peptide (KRRRALpSVASLPGL; SEQ ID NO:19) in the presence or absence of 0.2 μ M hymenialdisine or 1 μ M purvalanol B, compounds that are known or suspected inhibitors of CKI (Meijer, L. et al., 2000, *Chem. Biol.* 7:51-63; Knockaert et al., 2000, *supra*). This synthetic peptide substrate was chosen rather than casein because it is a highly selective substrate for mammalian CKI as a result of the preference this enzyme displays for phosphorylating serine residues immediately distal to serines that have previously been

modified by cAMP-dependent protein kinase ("PKA") (Flotow, H. et al., 1990, *J. Biol. Chem.* 265:14264-14269). Samples of fractions were also blotted and probed with antisera against a conserved internal epitope of TgCKI α which recognized both α and β forms of FLAGTgCKI (Figure 9B). A prominent peak of hymenialdisine and purvalanol B sensitive phosphopeptide kinase activity was identified in fractions 5-9 that correlated with the presence of a CKI-specific 38 kDa protein in the Western blot. A second less active peak was detected in fractions 14 and 15, which exhibited partial sensitivity to hymenialdisine and purvalanol B. Although a 60 kDa Western blot positive band detected with CKI antisera (Figure 9B) was found in these fractions, its presence did not correlate with phosphopeptide kinase activity. Furthermore, since replica blots probed with antisera selective for TgCKI β failed to detect the presence of cross-reacting protein in any of these fractions, it was concluded that this 60 kDa band is not TgCKI β . However, the phosphopeptide kinase activity of fractions 14 and 15 did correlate with the presence of *T. gondii* calmodulin domain-like protein kinase 1 (TgCDPK1), an enzyme that can also use casein as a substrate *in vitro* (not shown). Kinase assays with synthetic peptides that are preferred by TgCDPK1 (Syntide-2 and CaM kinase II substrate (281-291); Kieschnick et al., 2001, *supra*) confirmed a prominent peak of activity spanning these two fractions (not shown).

HiTrapQ fraction 7, which contained the peak activities for phosphopeptide kinase- and 38 kDa CKI protein, was subjected to hydrophobic interaction chromatography to further purify and concentrate the native CKI activity (Figure 9C). A peak of phosphopeptide kinase activity was eluted with a descending ammonium sulfate gradient which matched the presence of a 38 kDa band that was recognized by antisera specific for the TgCKI α isoform. (Figure 9D, 'CKI α -Ct'). The co-purification of phosphopeptide kinase activity with a 38 kDa TgCKI α protein through two column chromatography procedures suggested that the activity represents that of the native TgCKI α isoform. Fractions 14 and 15, containing the highest levels of this activity, were combined for further analysis.

A characterization of the enzymatic activities of preparations of recombinant TgCKI α or β and partially purified native TgCKI α isoforms is shown in Table I and Figure 10. A determination of the selectivities of the CKI enzyme preparations for casein and synthetic peptide substrates revealed qualitatively similar apparent K_m and V_{max} values. All of the CKI enzyme preparations yielded maximal activity with the phosphopeptide substrate. In the case of the recombinant TgCKI enzymes, where the greater degree of purity achieved permitted the determination of apparent V_{max} values, 2-3 fold higher levels of activity were obtained with this peptide than with either α - or β - casein. This similar increase was also seen in a comparison of maximal activity levels observed for the native TgCKI α preparation with these substrates (not normalized for protein, Figure 10B). Most non-phosphorylated synthetic peptides tested,

including those recognized by TgCDPK1 (Syntide-2 or CaM kinase II substrate), PKA (kemptide), some CKI enzymes [DDDEESITRR-OH (SEQ ID NO:17) or RRKDLHDDEEDEAMSITA-OH (SEQ ID NO:13)], and CKII [RRADDSDDDDD-OH (SEQ ID NO:14) or RRREEETEEE-OH (SEQ ID NO:20)] failed to show concentration dependent kinase activity. The single exception, a CKI peptide substrate (RRKDLHDDEEDEAMSITA-OH (SEQ ID NO:13)) yielded low levels of activity with $\text{FLAG-TgCKI}\beta$, but the activity was closer to background levels with native TgCKI α or $\text{FLAG-TgCKI}\beta$ enzymes (Table I, Figure 10C). Substrate selectivity and biochemical properties of the native EtCKI α enzyme were found to be very similar to the native TgCKI α . For example, in comparing the α -casein protein substrate with the preferred CKI phosphopeptide substrate, the phosphopeptide yielded a much higher V_{\max} (2.5X greater activity) than the α -casein substrate, similar to that seen with the recombinant TgCKI α and - β and native TgCKI α enzymes,

While purifying native *E. tenella* CKI α from unsporulated oocysts, a third and final cation exchange chromatography purification step resulted in active fractions of sufficient purity to identify a discrete ~40 kDa band on a silver stained SDS-PAGE gel (Figure 11A), which cross-reacted with CKI antisera and which correlated with CKI activity (Figure 11B). The activity was sensitive to a PKG inhibitor, Compound 20 (structure disclosed in PCT International Application PCT/US02/19507; International publication number WO 03/000682), with 100% inhibition at 250 nM.

To confirm the identity of native *E. tenella* CKI α , fractions 18, 19, and 20 from a EtCKI α purification were separated with SDS-PAGE. The gel was stained with Colloidal Coomassie, as well as silver stain (Figure 12). The ~40 kDa band was excised and digested with trypsin, and the resulting peptides were analyzed by nanoflow HPLC-micro-electrospray ionization tandem mass spectrometry. All the collected tandem mass spectra were processed using TurboSEQUENT to search against the *E. tenella* CKI α open reading frame. To confirm the identity of the tryptic peptides, an aliquot of the peptides was dried down and esterified in 2N methanolic HCl, which results in methyl groups (or 14 Da) being added to the acidic groups of every peptide. The resulting methyl esters were analyzed and processed in the same manner as described above. Molecular profiling showed conclusively that the LC-MS/MS profile of the ~40 kDa native protein matches the sequence of the cloned EtCKI α enzyme. Seven tryptic peptides were positively identified: SRHPQLIYESK (SEQ ID NO:37); TVLMLADQMLNR (SEQ ID NO:38); DIKPDNFLIGR (SEQ ID NO:39); TQSHIPYR (SEQ ID NO:40); YASVNTHLGIEQSR (SEQ ID NO:41); FEDRPDYSYLR (SEQ ID NO:42); and, DLFFR (SEQ ID NO:43) (represented by the shaded boxes in Figure 13). Five of these peptides were confirmed following methyl ester derivitization: SRHPQLIYESK (SEQ ID NO:37);

DIKPDNFLIGR (SEQ ID NO:39); YASVNTHLGIEQSR (SEQ ID NO:41); FEDRPDYSYLR (SEQ ID NO:42); and, DLFFR (SEQ ID NO:43).

Table 1. Kinetic parameters of TgCKI α and TgCKI β isoforms. Values for $V_{max(app)}$ (nmole min⁻¹ mg⁻¹) and $K_{m(app)}$ (μ M) were determined from titrations performed in a representative experiment (n=2 to 3). Catalytic parameters were measured in the presence of 2 μ M ATP and 200 μ M peptide or casein substrate. Parameters were calculated using curve-fitting software (Graphpad Prism) with "goodness of fit" values of $R^2 > 0.990 \pm 0.002$ for all values presented. Examples of plots from which these values were calculated are shown in Figure 10. 'NA' is not active.

Substrate	FLAGTgCKI β		FLAGTgCKI α		native TgCKI α
	$V_{max(app)}$	$K_{m(app)}$	$V_{max(app)}$	$K_{m(app)}$	$K_{m(app)}$
α -casein	12	5	8	4	3
β -casein	16	12	8	30	13
CKI peptide	8	15	NA	NA	NA
CKI phosphopeptide	37	79	26	80	20

EXAMPLE 4

Inhibitor Studies

Compounds - A collection of ATP-site competitive inhibitors of CDKs were used (Knockaert et al., 2002, *Trends Pharmacol. Sci.*). All but aminopurvalanol are currently available commercially (e.g., Alexis Corporation, Tocris, Calbiochem, and A.G. Scientific). They were stored as 10 mM stock solutions in DMSO.

Purification of recombinant TgCDPK1 and TgPKG - The TgCDPK1 open reading frame was amplified by PCR from a *Toxoplasma* cDNA library, using primers based on the published sequence (Kieschnick et al., 2001, *supra*): sense primer, AAAATGGGGCAGCAGGAAAGCACTCTTGCG (SEQ ID NO:44); and, antisense primer, GTTCCGCAGAGCTTCAAGAGCATCTGTT (SEQ ID NO:45). A C-terminal FLAG epitope tag was appended to the TgCDPK1 open reading frame by an additional round of PCR amplification (KOD polymerase) and the resulting DNA fragment was sub-cloned into a tubulin promoter expression vector (Donald and Liberator, 2002, *supra*). To facilitate the selection of stable transgenic lines, an HXGPRT minigene fragment was sliced into a unique expression-vector BamHI site, permitting mycophenolic acid selection in the parasite HXGPRT knock-out strain (Donald, R.G.K. et al., 1996, *J. Biol. Chem.* 271:14010-14019). Recombinant

FLAG-TgCDPK1 and FLAG-PKG enzymes were recovered from transgenic parasites by single step FLAG-affinity purification as described previously (Donald and Liberator, 2002, *supra*). The activity of the FLAG-TgCDPK1 enzyme closely resembled properties of the native TgCDPK1 enzyme with respect to sensitivity to calcium and apparent K_m and V_{max} values (Kieschnick et al, 2001, *supra*).

Results - The sensitivity of CKI enzymes and *T. gondii* parasites to a variety of antimetabolic CDK inhibitors was evaluated, and the results are shown in Table 2. Compound 1, a selective inhibitor of coccidian parasite PKG with well-characterized anti-parasitic properties (Nare, B. et al., 2002, *supra*; Gurnett, et al., 2002, *supra*; Donald et al., 2002, *supra*; US Pat. No. 6,291,480), was included as an additional reference compound. The first five CDK compounds listed (Compounds 2-6) belong to the 2,6,9-trisubstituted purine structural class of which the purvalanols are members (Meijer and Raymond, 2003, *Accounts Chem. Res.* 36:417-425). Aminopurvalanol, purvalanol A and purvalanol B are highly potent and selective CDK inhibitors with relatively low activity against other kinases in mammalian cells (Chang, Y.T. et al., 1999, *Chem. Biol.* 6:361-375; Gray, N.S. et al., 1998, *Science* 281:533-538; Knockaert et al., 2000, *supra*). The structures of these compounds are shown in Figure 14. Indirubin-3'-monoxime, an active ingredient of a Chinese anti-leukemic herbal remedy, is a less selective compound with activity against a number of other mammalian kinases including glycogen synthase kinase (GSK-3), AMP-activated protein kinase (AMPK), a tyrosine kinase (LCK), and serum activated protein kinase (SGK) (Leclerc, S. et al., 2001, *J. Biol. Chem.* 276:251-260; Damiens, E. et al., 2001, *Oncogene* 20:3786-3797; Marko, D. et al., 2001, *Br. J. Cancer* 84:283-289; Bain, J. et al., 2003, *Biochem. J.* 371:199-204). Hymenialdisine, a marine sponge constituent, is a potent inhibitor of human CDKs, but also of GSK-3 and CKI (Meijer, L. et al., 2000, *Chem. Biol.* 7:51-63). Kenpaullone and alsterpaullone, representatives of the paullone family of compounds, display good selectivity toward CDKs, although they are also potent inhibitors of GSK-3 (Zaharevitz, D.W. et al., 1999, *Cancer Res.* 59:2566-2569; Knockaert, M. et al., 2002, *J. Biol. Chem.* 277:25493-25501; Bain et al., 2003, *supra*). To broaden the scope of this enzyme-target selectivity study, the activity of these compounds against *T. gondii* PKG and CDPK1 was also assessed.

Two of the purvalanols, purvalanol B and aminopurvalanol, showed excellent activity against *T. gondii* in the uracil uptake assay, a parasite-specific metabolic assay performed with infected HFF cell monolayers (Pfefferkorn and Pfefferkorn, 1977, *supra*). Other members of this structural class showed only mediocre activity (purvalanol A, roscovitine) or no activity (olomoucine) against cultured parasites. The 260 nM and 375 nM IC_{50} values obtained respectively for purvalanol B and aminopurvalanol are similar to those observed with Compound

1 (IC₅₀ 200 nM). These same compounds, but not purvalanol A, roscovitine or olomoucine, also showed good activity against recombinant or native forms of TgCKI α (IC₅₀ 40-60 nM). The only other CDK inhibitor with sub- μ M activity against *T. gondii* was indirubin-3'-monoxime, which gave an IC₅₀ value of 500 nM. This compound showed modest activity against TgCKI α (IC₅₀ 172 nM), but also against the CDPK1 enzyme (IC₅₀ 175 nM). Hymenialdisine showed no antiparasitic activity, but was a potent inhibitor of both α and β isoforms of TgCKI (IC₅₀ 7-10 nM) as well as rodent CKI δ (IC₅₀ 60 nM). Neither kenpaullone nor alsterpaullone showed any activity against the parasite kinases tested. Compared with kenpaullone, which was inactive in the whole cell assay, alsterpaullone showed some antiparasitic activity at low micromolar levels (IC₅₀ 1.2 μ M). Alsterpaullone is at least 10-fold more potent against mammalian CDK1 or CDK5 than kenpaullone (Knockaert, M. et al, 2002, *J. Biol. Chem.*). Native EtCKI α shows similar sensitivities as compared to TgCKI enzymes to these reference compounds (Table 2).

Biochemical characterization of the partially purified native EtCKI α isoform (e.g., fraction 19, Figure 13) shows that it resembles its *T. gondii* counterpart with respect to substrate selectivity and sensitivity to reference compounds (CDK inhibitors) and imidazopyridines (PKG inhibitor leads). Table 3 summarizes the sensitivity of various coccidian protein kinases to two imidazopyridine compounds, Compound 20 (PCT International Application PCT/US02/19507; International publication no. WO 03/000682) and Compound 39 (see Figure 15). *T. gondii* and *E. tenella* CKI enzymes show modest levels of sensitivity to these compounds.

Table 2. Sensitivity of *T. gondii* PKG, CKI α and $-\beta$ (expressed and native), and CDPK1; and native *E. tenella* CKI α , and rat CKI δ to inhibitors of mammalian CDK. PKG inhibitor Compound 1 was included as a control. Enzyme- and whole cell- metabolic activities were assayed as described above in the presence of increasing concentrations of compound. IC₅₀ values (nM) were calculated from dose response curves. Submicromolar values are highlighted in grey. 'NT' is not tested. Structures of purvalanols A, B and aminopurvalanol are shown in Figure 14. The errors shown for some values represent standard deviations resulting from additional titration experiments (n=2-3).

	<i>Inhibitor</i>	³ [H]Uracil	<i>TgPKG</i>	<i>TgCDPK1</i>	<i>TgCKIβ</i>	<i>TgCKIα</i>	native	native	Rat
		<i>Uptake</i>					<i>TgCKIα</i>	<i>EtCKIα</i>	<i>CKIδ</i>
5	1. Compound 1	200 ± 100	0.7 ± 0.3	63 ± 2	4,200	107 ± 19	95 ± 28	350	2,230
	2. Purvalanol A	1,230	3,500	4140	>5,000	>5,000	>5,000	4,900	>5,000
	3. Purvalanol B	260 ± 84	1,900	770 ± 200	606	38 ± 12	63	94	3,600
	4. Aminopurvalanol	375 ± 21	2,200	730 ± 200	3760	42 ± 7	61	98	3,500
	5. Roscovitine	>5,000	10,000	>5,000	>5,000	1,200	1,200	1,700	>5,000
10	6. Olomoucine	>5,000	10,000	4200	>5,000	>5,000	>5,000	NT	>5,000
	7. Indirubin-3'-monoxime	500 ± 100	1865	172 ± 4	1340	124 ± 21	550	NT	>5,000
	8. Hymenialdisine	>10,000	2136	175 ± 33	11 ± 6	7 ± 3	10	NT	60
	9. Alsterpauillone	1,220	>5,000	>5,000	>5,000	>5,000	NT	NT	2,300
15	10. Kenpauillone	>5,000	>5,000	>5,000	>5,000	>5,000	NT	NT	>5,000

Table 3. Coccidian CKIα and CDPK1 enzymes are secondary targets of lead PKG inhibitors. Sensitivity of recombinant- or partially purified native enzymes to PKG inhibitors was determined from dose response titrations (nM IC₅₀ values). Recombinant enzymes were purified from *E. coli* and native enzymes from extracts of *Toxoplasma* tachyzoites (TgCKIα) or *Eimeria* oocysts (EtCKIα and EtPKG).

	<i>Enzyme</i>	<i>Compound 20</i>	<i>Compound 39</i>
	FLAGTgCDPK1	1.1	0.5
25	FLAGTgCDPK3	>10,000	>10,000
	FLAGEtCDPK1	413	126
	FLAGEtCDPK2	>10,000	>10,000
	nativeTgCKIα	18	15
	nativeEtCKIα	84	69
30	nativeEtPKG	0.7	0.5